FILE 'HOME' ENTERED AT 14:59:34 ON 19 SEP 2003

=> fil .bec

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

FILES 'MEDLINE, SCISEARCH, LIFESCI, BIOTECHDS, BIOSIS, EMBASE, HCAPLUS, NTIS, ESBIOBASE, BIOTECHNO, WPIDS' ENTERED AT 14:59:45 ON 19 SEP 2003 ALL COPYRIGHTS AND RESTRICTIONS APPLY. SEE HELP USAGETERMS FOR DETAILS.

11 FILES IN THE FILE LIST

=> s alpha amylase#

FILE 'MEDLINE'

454272 ALPHA

20114 AMYLASE#

L1 4463 ALPHA AMYLASE#

(ALPHA(W)AMYLASE#)

FILE 'SCISEARCH'

642997 ALPHA

16176 AMYLASE#

L2 7215 ALPHA AMYLASE#

(ALPHA (W) AMYLASE#)

FILE 'LIFESCI'

147374 "ALPHA"

4283 AMYLASE#

L3 2572 ALPHA AMYLASE#

("ALPHA" (W) AMYLASE#)

FILE 'BIOTECHDS'

23979 ALPHA

4870 AMYLASE#

L4 3195 ALPHA AMYLASE#

(ALPHA(W)AMYLASE#)

FILE 'BIOSIS'

596697 ALPHA

26984 AMYLASE#

L5 9606 ALPHA AMYLASE#

(ALPHA (W) AMYLASE#)

FILE 'EMBASE'

508813 "ALPHA"

14781 AMYLASE#

L6 3284 ALPHA AMYLASE#

("ALPHA"(W)AMYLASE#)

FILE 'HCAPLUS'

1434449 ALPHA

42789 AMYLASE#

L7 17646 ALPHA AMYLASE#

(ALPHA(W)AMYLASE#)

FILE 'NTIS'

28426 ALPHA

163 AMYLASE#

L8 60 ALPHA AMYLASE#

(ALPHA (W) AMYLASE#)

FILE 'ESBIOBASE'

176805 ALPHA

3749 AMYLASE#

L9

1783 ALPHA AMYLASE#

(ALPHA(W)AMYLASE#)

FILE 'BIOTECHNO'

183303 ALPHA

4113 AMYLASE#

L10 2087 ALPHA AMYLASE#

(ALPHA(W)AMYLASE#)

FILE 'WPIDS'

164909 ALPHA

5095 AMYLASE#

L11 2102 ALPHA AMYLASE#

(ALPHA(W)AMYLASE#)

TOTAL FOR ALL FILES

L12 54013 ALPHA AMYLASE#

=> s 112(5a)gene/q

FILE 'MEDLINE'

L13 568 L1 (5A)GENE/Q

FILE 'SCISEARCH'

L14 808 L2 (5A)GENE/Q

FILE 'LIFESCI'

L15 578 L3 (5A)GENE/Q

FILE 'BIOTECHDS'

L16 732 L4 (5A)GENE/Q

FILE 'BIOSIS'

L17 1003 L5 (5A)GENE/Q

FILE 'EMBASE'

L18 446 L6 (5A)GENE/Q

FILE 'HCAPLUS'

L19 1713 L7 (5A)GENE/Q

FILE 'NTIS'

L20 5 L8 (5A)GENE/Q

FILE 'ESBIOBASE'

L21 273 L9 (5A)GENE/Q

FILE 'BIOTECHNO'

L22 495 L10(5A)GENE/Q

FILE 'WPIDS'

L23 156 L11(5A)GENE/Q

TOTAL FOR ALL FILES

L24 6777 L12(5A) GENE/Q

=> s hyperthermophil? or thermophil?

FILE 'MEDLINE'

1419 HYPERTHERMOPHIL?

7477 THERMOPHIL?

L25 8631 HYPERTHERMOPHIL? OR THERMOPHIL?

FILE 'SCISEARCH'

2090 HYPERTHERMOPHIL?

12850 THERMOPHIL?

L26 14345 HYPERTHERMOPHIL? OR THERMOPHIL?

FILE 'LIFESCI'

1155 HYPERTHERMOPHIL?

7671 THERMOPHIL?

L27 8294 HYPERTHERMOPHIL? OR THERMOPHIL?

FILE 'BIOTECHDS'

282 HYPERTHERMOPHIL?

5266 THERMOPHIL?

L28 5325 HYPERTHERMOPHIL? OR THERMOPHIL?

FILE 'BIOSIS'

1834 HYPERTHERMOPHIL?

16462 THERMOPHIL?

L29 17388 HYPERTHERMOPHIL? OR THERMOPHIL?

FILE 'EMBASE'

1374 HYPERTHERMOPHIL?

8071 THERMOPHIL?

L30 8712 HYPERTHERMOPHIL? OR THERMOPHIL?

FILE 'HCAPLUS'

1988 HYPERTHERMOPHIL?

17169 THERMOPHIL?

L31 18713 HYPERTHERMOPHIL? OR THERMOPHIL?

FILE 'NTIS'

31 HYPERTHERMOPHIL?

485 THERMOPHIL?

L32 505 HYPERTHERMOPHIL? OR THERMOPHIL?

FILE 'ESBIOBASE'

1267 HYPERTHERMOPHIL?

4548 THERMOPHIL?

L33 5543 HYPERTHERMOPHIL? OR THERMOPHIL?

FILE 'BIOTECHNO'

1252 HYPERTHERMOPHIL?

6720 THERMOPHIL?

L34 7311 HYPERTHERMOPHIL? OR THERMOPHIL?

FILE 'WPIDS'

47 HYPERTHERMOPHIL?

1931 THERMOPHIL?

L35 1956 HYPERTHERMOPHIL? OR THERMOPHIL?

TOTAL FOR ALL FILES

L36 96723 HYPERTHERMOPHIL? OR THERMOPHIL?

=> s 124 and 136

FILE 'MEDLINE'

L37 29 L13 AND L25

FILE 'SCISEARCH'

L38 36 L14 AND L26

FILE 'LIFESCI'

L39 37 L15 AND L27

FILE 'BIOTECHDS'

L40 94 L16 AND L28

FILE 'BIOSIS'

L41 46 L17 AND L29

FILE 'EMBASE'

L42 39 L18 AND L30

FILE 'HCAPLUS'

L43 67 L19 AND L31

FILE 'NTIS'

L44 1 L20 AND L32

FILE 'ESBIOBASE'

L45 25 L21 AND L33

FILE 'BIOTECHNO'

L46 40 L22 AND L34

FILE 'WPIDS'

L47 8 L23 AND L35

TOTAL FOR ALL FILES

L48 422 L24 AND L36

=> s 148 not 2002-2003/py

FILE 'MEDLINE'

897062 2002-2003/PY

L49 26 L37 NOT 2002-2003/PY

FILE 'SCISEARCH'

1610344 2002-2003/PY

L50 33 L38 NOT 2002-2003/PY

FILE 'LIFESCI'

137782 2002-2003/PY

L51 32 L39 NOT 2002-2003/PY

FILE 'BIOTECHDS'

36013 2002-2003/PY

L52 90 L40 NOT 2002-2003/PY

FILE 'BIOSIS'

812724 2002-2003/PY

L53 40 L41 NOT 2002-2003/PY

FILE 'EMBASE'

739028 2002-2003/PY

L54 34 L42 NOT 2002-2003/PY

FILE 'HCAPLUS'

1760625 2002-2003/PY

L55 58 L43 NOT 2002-2003/PY

FILE 'NTIS'

17588 2002-2003/PY

L56 1 L44 NOT 2002-2003/PY

FILE 'ESBIOBASE'

467486 2002-2003/PY

L57 21 L45 NOT 2002-2003/PY

FILE 'BIOTECHNO'

203275 2002-2003/PY

L58 34 L46 NOT 2002-2003/PY

FILE 'WPIDS'

1725290 2002-2003/PY

L59 6 L47 NOT 2002-2003/PY

TOTAL FOR ALL FILES

L60 375 L48 NOT 2002-2003/PY

=> dup rem 160

PROCESSING COMPLETED FOR L60

L61 134 DUP REM L60 (241 DUPLICATES REMOVED)

=> d tot

L61 ANSWER 1 OF 134 HCAPLUS COPYRIGHT 2003 ACS on STN

Novel, thermostable family-13-like glycoside hydrolase from Methanococcus jannaschii

SO Folia Microbiologica (Prague, Czech Republic) (2001), 46(6), 475-481 CODEN: FOMIAZ; ISSN: 0015-5632

AU Kim, J.-W.; Terc, H. A.; Flowers, L. O.; Whiteley, M.; Peeples, T. L.

AN 2002:140158 HCAPLUS

DN 137:59380

ΑU

L61 ANSWER 2 OF 134 MEDLINE on STN DUPLICATE 1

TI Biochemical confirmation and characterization of the family-57-like alpha-amylase of Methanococcus jannaschii.

SO FOLIA MICROBIOLOGICA, (2001) 46 (6) 467-73. Journal code: 0376757. ISSN: 0015-5632.

AU Kim J W; Flowers L O; Whiteley M; Peeples T L

AN 2002167588 MEDLINE

L61 ANSWER 3 OF 134 MEDLINE on STN DUPLICATE 2

Novel glucoamylase-type enzymes from Thermoactinomyces vulgaris and Methanococcus jannaschii whose genes are found in the flanking region of the alpha-amylase genes.

SO APPLIED MICROBIOLOGY AND BIOTECHNOLOGY, (2001 Aug) 56 (3-4) 465-73. Journal code: 8406612. ISSN: 0175-7598.

Uotsu-Tomita R; Tonozuka T; Sakai H; Sakano Y

AN 2001499375 MEDLINE

L61 ANSWER 4 OF 134 MEDLINE on STN DUPLICATE 3

TI Cloning and expression of alpha-amylase from the hyperthermophilic archaeon Pyrococcus woesei in the moderately halophilic bacterium Halomonas elongata.

SO JOURNAL OF APPLIED MICROBIOLOGY, (2000 Mar) 88 (3) 495-503. Journal code: 9706280. ISSN: 1364-5072.

AU Frillingos S; Linden A; Niehaus F; Vargas C; Nieto J J; Ventosa A; Antranikian G; Drainas C

AN 2000212011 MEDLINE

L61 ANSWER 5 OF 134 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN

TI Cloning and expression of the **gene** encoding novel **alpha** -amylase from Sulfolobus shibatae in E. coli.

SO Weishengwu Xuebao, (June, 2000) Vol. 40, No. 3, pp. 323-326. print. ISSN: 0001-6209.

AU Liu Li (1); Chen Wei (1); Jin Cheng

AN 2001:49285 BIOSIS

L61 ANSWER 6 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN

TI Characterization of hygromycin-resistant transformants of

thermophilic fungus Thermomyces lanuginosus;

plasmid pMP6-mediated hygromycin-resistance gene transfer

together with alpha-amylase, glucoamylase, polygalacturonase and endo-1,4-beta-D-xylanase production

World J.Microbiol.Biotechnol.; (2000) 16, 3, 303-06 SO

ISSN: 0959-3993 CODEN: WJMBEY

Chadha B S; Kaur R; Saini H S; Singh S ΑU

2000-10928 BIOTECHDS AN

DUPLICATE 4 MEDLINE on STN ANSWER 7 OF 134 L61

Single-step purification of a recombinant thermostable alpha-amylase after TIsolubilization of the enzyme from insoluble aggregates.

JOURNAL OF CHROMATOGRAPHY. B, BIOMEDICAL SCIENCES AND APPLICATIONS, (2000 SO Jan 14) 737 (1-2) 253-9. Journal code: 9714109. ISSN: 1387-2273.

Linden A; Niehaus F; Antranikian G ΑU

MEDLINE 2000143297 AN

L61 ANSWER 8 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN

Releasing profiles of gene products from recombinant Escherichia coli in a high-voltage pulsed electric field

BIOCHEMICAL ENGINEERING JOURNAL, (JUN 2000) Vol. 5, No. 2, pp. 149-155. Publisher: ELSEVIER SCIENCE SA, PO BOX 564, 1001 LAUSANNE, SWITZERLAND. SO ISSN: 1369-703X.

Ohshima T (Reprint); Hama Y; Sato M ΑU

2000:414547 SCISEARCH AN

DUPLICATE 5 MEDLINE on STN L61 ANSWER 9 OF 134

Cloning and expression of an alpha-amylase encoding TIgene from the hyperthermophilic archaebacterium Thermococcus hydrothermalis and biochemical characterisation of the recombinant enzyme.

FEMS MICROBIOLOGY LETTERS, (2000 May 1) 186 (1) 67-71. SO Journal code: 7705721. ISSN: 0378-1097.

Leveque E; Haye B; Belarbi A AU

MEDLINE 2000395702 AN

DUPLICATE 6 MEDLINE on STN L61 ANSWER 10 OF 134

Engineering direct fructose production in processed potato tubers by expressing a bifunctional alpha-amylase/glucose isomerase gene complex.

BIOTECHNOLOGY AND BIOENGINEERING, (2000 Oct 5) 70 (1) 9-16. SO Journal code: 7502021. ISSN: 0006-3592.

Beaujean A; Ducrocq-Assaf C; Sangwan R S; Lilius G; Bulow L; ΑU Sangwan-Norreel B S

MEDLINE 2001028442 AN

ANSWER 11 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN

L61 Production of recombinant thermophilic alpha-amylase activity TI at low pH;

Thermococcus hydrothermalis recombinant enzyme production via plasmid pEAMY101 and plasmid p662EL100 expression in Escherichia coli

Leveque E; Belarbi A; Haye B ΑU

2000-03337 BIOTECHDS ΑN

FR 2778412 12 Nov 1999 PΙ

L61 ANSWER 12 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN

A unique chitinase with dual active sites and triple substrate binding sites from the hyperthermophilic archaeon Pyrococcus

kodakaraensis KOD1 APPLIED AND ENVIRONMENTAL MICROBIOLOGY, (DEC 1999) Vol. 65, No. 12, pp. SO

5338-5344. Publisher: AMER SOC MICROBIOLOGY, 1325 MASSACHUSETTS AVENUE, NW, WASHINGTON, DC 20005-4171.

ISSN: 0099-2240.

Tanaka T; Fujiwara S; Nishikori S; Fukui T; Takagi M; Imanaka T (Reprint) ΑU

- AN 1999:949061 SCISEARCH
- L61 ANSWER 13 OF 134 HCAPLUS COPYRIGHT 2003 ACS on STN
- TI Coordinate transcriptional control in the hyperthermophilic archaeon Sulfolobus solfataricus
- SO Journal of Bacteriology (1999), 181(13), 3920-3927 CODEN: JOBAAY; ISSN: 0021-9193
- AU Haseltine, Cynthia; Montalvo-Rodriguez, Rafael; Bini, Elisabetta; Carl, Audrey; Blum, Paul
- AN 1999:414107 HCAPLUS
- DN 131:195320
- L61 ANSWER 14 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN
- Purification and characterization of an extremely thermostable cyclomaltodextrin glucanotransferase from a newly isolated hyperthermophilic archaeon, a Thermococcus sp.
- SO APPLIED AND ENVIRONMENTAL MICROBIOLOGY, (MAY 1999) Vol. 65, No. 5, pp. 1991-1997.
 Publisher: AMER SOC MICROBIOLOGY, 1325 MASSACHUSETTS AVENUE, NW,

WASHINGTON, DC 20005-4171.

ISSN: 0099-2240.

- AU Tachibana Y (Reprint); Kuramura A; Shirasaka N; Suzuki Y; Yamamoto T; Fujiwara S; Takagi M; Imanaka T
- AN 1999:353855 SCISEARCH
- L61 ANSWER 15 OF 134 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
- TI Extragenic pleiotropic mutations that repress glycosyl hydrolase expression in the hyperthermophilic archaeon Sulfolobus solfataricus.
- SO Genetics, (Aug., 1999) Vol. 152, No. 4, pp. 1353-1361. ISSN: 0016-6731.
- AU Haseltine, Cynthia; Montalvo-Rodriguez, Rafael; Carl, Audrey; Bini, Elisabetta; Blum, Paul (1)
- AN 1999:416085 BIOSIS
- L61 ANSWER 16 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
- Fermentation of starch by Klebsiella oxytoca P2, containing plasmids with alpha-amylase and pullulanase genes;

for use in enzyme production or ethanol production SO Biotechnol.Bioeng.; (1999) 65, 6, 673-76

CODEN: BIBIAU ISSN: 0006-3592

- AU dos Santos V L; Fernandes Araujo E; Goncalves de Barros E; Vieira Guimaraes W
- AN 2000-00826 BIOTECHDS
- L61 ANSWER 17 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
- TI Production of alpha-amylase in fed-batch cultures of vgb+ and vbgrecombinant Escherichia coli: some observations;

Bacillus stearothermophilus recombinant enzyme production using host expressing (or not expressing) Vitreoscilla hemoglobin

- SO Biotechnol.Prog.; (1999) 15, 4, 640-45 CODEN: BIPRET ISSN: 8756-7938
- AU Enayati N; Tari C; *Parulekar S J; Stark B C; Webster D A
- AN 1999-11780 BIOTECHDS
- L61 ANSWER 18 OF 134 HCAPLUS COPYRIGHT 2003 ACS on STN
- TI Purification and characterization of the heat-labile .alpha.-amylase secreted by the psychrophilic bacterium TAC 240B
- SO Canadian Journal of Microbiology (1999), 45(6), 452-457 CODEN: CJMIAZ; ISSN: 0008-4166
- AU Chessa, Jean-Pierre; Feller, Georges; Gerday, Charles
- AN 1999:504065 HCAPLUS
- DN 131:254106

- L61 ANSWER 19 OF 134 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
- TI Close evolutionary relatedness of alpha-amylases from Archaea and plants.
- SO Journal of Molecular Evolution, (April, 1999) Vol. 48, No. 4, pp. 421-426. ISSN: 0022-2844.
- AU Janecek, Stefan (1); Leveque, Emmanuel; Belarbi, Abdel; Haye, Bernard
- AN 1999:220372 BIOSIS
- L61 ANSWER 20 OF 134 HCAPLUS COPYRIGHT 2003 ACS on STN
- TI Bioengineering of amylase and xylose isomerase thermozymes
- SO Special Publication Royal Society of Chemistry (1999), 246 (Recent Advances in Carbohydrate Bioengineering), 253-262 CODEN: SROCDO; ISSN: 0260-6291
- AU Zeikus, J. G.; Savchenko, A.; Sriprapundh, D.; Vieille, Claire
- AN 2000:29433 HCAPLUS
- DN 132:193269
- L61 ANSWER 21 OF 134 HCAPLUS COPYRIGHT 2003 ACS on STN
- TI Amylase and 16S rRNA genes from a hyperthermophilic archaebacterium
- SO Journal of Applied Microbiology (1999), 86(1), 93-107 CODEN: JAMIFK; ISSN: 1364-5072
- AU Jones, R. A.; Jermiin, L. S.; Easteal, S.; Patel, B. K. C.; Beacham, I. R.
- AN 1999:138511 HCAPLUS
- DN 131:40353
- L61 ANSWER 22 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
- TI Regulation of the expression of amy TO1 encoding a thermostable alpha-amylase from Streptomyces sp. TO1, in its original host and in Streptomyces lividans TK24;
 - thermostable alpha-amylase production
- SO FEMS Microbiol.Lett.; (1999) 181, 1, 31-39
- CODEN: FMLED7 ISSN: 0378-1097
- AU Mellouli L; Guerineau M; Bejar S; Virolle M J
- AN 2000-00399 BIOTECHDS
- L61 ANSWER 23 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
- TI Synthetic and excretion of alpha-amylase in vgb+ and vgb-recombinant Escherichia coli: a comparative study;
 - Bacillus stearothermophilus recombinant alpha-amylase production
- SO Biotechnol.Bioeng.; (1998) 59, 6, 673-78 CODEN: BIBIAU ISSN: 0006-3592
- AU Tari C; *Parulekar S J; Stark B C; Webster D A
- AN 1999-12957 BIOTECHDS
- L61 ANSWER 24 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN
- TI Sequence of archaeal Methanococcus jannaschii alphaamylase contains features of families 13 and 57 of glycosyl hydrolases: A trace of their common ancestor?
- FOLIA MICROBIOLOGICA, (FEB 1998) Vol. 43, No. 2, pp. 123-128.

 Publisher: FOLIA MICROBIOLOGICA, INST MICROBIOLOGY, VIDENSKA 1083, PRAGUE
 4, CZECH REPUBLIC 142 20.

 ISSN: 0015-5632.
- AU Janecek S (Reprint)
- AN 1998:208636 SCISEARCH
- L61 ANSWER 25 OF 134 HCAPLUS COPYRIGHT 2003 ACS on STN
- TI Extracellular .alpha.-amylase from Thermus filiformis Ork A2: purification and biochemical characterization
- SO Extremophiles (1998), 2(1), 23-32 CODEN: EXTRFI; ISSN: 1431-0651
- AU Egas, Maria C. V.; da Costa, Milton S.; Cowan, Don A.; Pires, Euclides M. V.
- AN 1998:144647 HCAPLUS
- DN 128:241074

- L61 ANSWER 26 OF 134 HCAPLUS COPYRIGHT 2003 ACS on STN
- Hyperthermostable extracellular alpha-amylase from pyrococcus furiosus ΤI
- Book of Abstracts, 216th ACS National Meeting, Boston, August 23-27 SO (1998), BTEC-019 Publisher: American Chemical Society, Washington, D. C. CODEN: 66KYA2
- Savchenko, A.; Dong, G.; Vieille, C.; Zeikus, G. J. ΑU
- 1998:528470 HCAPLUS AN
- L61 ANSWER 27 OF 134 MEDLINE on STN

DUPLICATE 8

- ΤI alpha-Amylase gene of thermophilic Streptomyces sp. TO1: nucleotide sequence, transcriptional and amino acid sequence analysis.
- FEMS MICROBIOLOGY LETTERS, (1998 Mar 1) 160 (1) 17-23. SO Journal code: 7705721. ISSN: 0378-1097.
- Mellouli L; Ghorbel R; Virolle M J; Bejar S ΑU
- 1998156111 MEDLINE ΑN
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- Isolation and analysis of genes for amylolytic enzymes of the TΙ hyperthermophilic bacterium Thermotoga maritima.
- FEMS MICROBIOLOGY LETTERS, (1998 Jan 1) 158 (1) 9-15. SO Journal code: 7705721. ISSN: 0378-1097.
- Bibel M; Brettl C; Gosslar U; Kriegshauser G; Liebl W ΑU
- MEDLINE AN1998115241
- ANSWER 29 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61
- Hyperthermostable extracellular alpha-amylase from Pyrococcus furiosus; TΤ thermophilic bacterium recombinant enzyme production and

characterization (conference abstract)

- SO Abstr.Pap.Am.Chem.Soc.; (1998) 216 Meet. Pt.3, BTEC019
 - ISSN: 0065-7727 CODEN: ACSRAL
 - 216th ACS National Meeting, Boston, MA, USA, 23-27 August, 1998, 216 Meet., Pt.3, 1998.
- Savchenko A; Dong G; Vieille C; Zeikus G J ΑU
- 1999-14174 BIOTECHDS AN
- ANSWER 30 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61
- Vector for secretion for use in lactic acid bacteria, and production of a ΤI protein by use of the vector;

recombinant enzyme e.g. amylase, peptidase or protease production

- 1997-13446 BIOTECHDS AN
- JP 09234078 9 Sep 1997 PΙ
- ANSWER 31 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61
- Termamyl-like alpha-amylase variants with improved properties; TIenzyme engineering and expression in Bacillus spp.
- Svensden A; Borchert T V; Bisgard-Frantzen H ΑU
- 1998-01800 BIOTECHDS ΑN
- PΙ WO 9741213 6 Nov 1997
- L61 ANSWER 32 OF 134 HCAPLUS COPYRIGHT 2003 ACS on STN
- Cloning and expression of the gene encoding super-thermostable . TIalpha. - amylase the hyperthermophilic
 - Pyrococcus strain KOD-1, and characterization and use of the enzyme
- Jpn. Kokai Tokkyo Koho, 12 pp. SO
 - CODEN: JKXXAF
- Imanaka, Tadayuki; Tachibana, Yoshinaga; Suzuki, Yuji; Kojima, Iwao; IN Utsura, Kensaku
- 1997:470120 HCAPLUS AN
- DN 127:77927
 - APPLICATION NO. DATE PATENT NO. KIND DATE _____
- JP 09173077 A2 19970708 JP 1996-191138 19960719 PΙ

- L61 ANSWER 33 OF 134 MEDLINE on STN DUPLICATE 11
- TI Cloning, sequencing, characterization, and expression of an extracellular alpha-amylase from the **hyperthermophilic** archaeon Pyrococcus furiosus in Escherichia coli and Bacillus subtilis.
- SO JOURNAL OF BIOLOGICAL CHEMISTRY, (1997 Jun 27) 272 (26) 16335-42. Journal code: 2985121R. ISSN: 0021-9258.
- AU Jorgensen S; Vorgias C E; Antranikian G
- AN 97341170 MEDLINE
- L61 ANSWER 34 OF 134 MEDLINE on STN DUPLICATE 12
- TI Application of the extracellular alpha-amylase gene from Streptococcus bovis 148 to construction of a secretion vector for yogurt starter strains.
- SO APPLIED AND ENVIRONMENTAL MICROBIOLOGY, (1997 Nov) 63 (11) 4593-6. Journal code: 7605801. ISSN: 0099-2240.
- AU Satoh E; Ito Y; Sasaki Y; Sasaki T
- AN 1998027396 MEDLINE
- L61 ANSWER 35 OF 134 MEDLINE on STN DUPLICATE 13
- TI Cloning, sequencing, and expression of the **gene** encoding extracellular **alpha-amylase** from Pyrococcus furiosus and biochemical characterization of the recombinant enzyme.
- SO APPLIED AND ENVIRONMENTAL MICROBIOLOGY, (1997 Sep) 63 (9) 3569-76. Journal code: 7605801. ISSN: 0099-2240.
- AU Dong G; Vieille C; Savchenko A; Zeikus J G
- AN 97438520 MEDLINE
- L61 ANSWER 36 OF 134 MEDLINE on STN DUPLICATE 14
- TI Characterization of the gene encoding an extracellular laccase of Myceliophthora thermophila and analysis of the recombinant enzyme expressed in Aspergillus oryzae.
- SO APPLIED AND ENVIRONMENTAL MICROBIOLOGY, (1997 Aug) 63 (8) 3151-7. Journal code: 7605801. ISSN: 0099-2240.
- AU Berka R M; Schneider P; Golightly E J; Brown S H; Madden M; Brown K M; Halkier T; Mondorf K; Xu F
- AN 97394941 MEDLINE
- L61 ANSWER 37 OF 134 MEDLINE ON STN DUPLICATE 15
- Properties and **gene** structure of the Thermotoga maritima **alpha-amylase** AmyA, a putative lipoprotein of a **hyperthermophilic** bacterium.
- SO JOURNAL OF BACTERIOLOGY, (1997 Feb) 179 (3) 941-8. Journal code: 2985120R. ISSN: 0021-9193.
- AU Liebl W; Stemplinger I; Ruile P
- AN 97158692 MEDLINE
- L61 ANSWER 38 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN DUPLICATE 16
- TI Study of stability of recombinant plasmids during the continuous culture of Bacillus stearothermophilus NUB3621 in nonselective medium
- SO BIOTECHNOLOGY AND BIOENGINEERING, (5 MAR 1997) Vol. 53, No. 5, pp. 507-514.
 - Publisher: JOHN WILEY & SONS INC, 605 THIRD AVE, NEW YORK, NY 10158-0012. ISSN: 0006-3592.
- AU Brigidi P (Reprint); GonzalezVara A; Rossi M; Matteuzzi D
- AN 97:193236 SCISEARCH
- L61 ANSWER 39 OF 134 LIFESCI COPYRIGHT 2003 CSA on STN DUPLICATE 17
- TI Study of stability of recombinant plasmids during the continuous culture of Bacillus stearothermophilus NUB3621 in nonselective medium
- SO BIOTECHNOL. BIOENG., (1997) vol. 53, no. 5, pp. 507-514. ISSN: 0006-3952.
- AU Brigidi, P.; Gonzalez-Vara, A.; Rossi, M.; Matteuzzi, D.

- L61 ANSWER 40 OF 134 MEDLINE on STN DUPLICATE 18
- TI A gene encoding for an alpha-amylase from thermophilic Bacillus sp. strain TS-23 and its expression in Escherichia coli.
- SO JOURNAL OF APPLIED MICROBIOLOGY, (1997 Mar) 82 (3) 325-34. Journal code: 9706280. ISSN: 1364-5072.
- AU Lin L L; Hsu W H; Chu W S
- AN 2002694119 MEDLINE
- L61 ANSWER 41 OF 134 MEDLINE on STN DUPLICATE 19
- TI Gene cloning and expression of new trehalose-producing enzymes from the hyperthermophilic archaeum Sulfolobus solfataricus KM1.
- SO BIOSCIENCE, BIOTECHNOLOGY, AND BIOCHEMISTRY, (1996 Nov) 60 (11) 1882-5. Journal code: 9205717. ISSN: 0916-8451.
- AU Kobayashi K; Kato M; Miura Y; Kettoku M; Komeda T; Iwamatsu A
- AN 97141610 MEDLINE
- L61 ANSWER 42 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
- TI Gene analysis of trehalose-producing enzymes from hyperthermophilic archaea in Sulfolobales;
- hyperthermophilic bacterium
- SO Biosci.Biotechnol.Biochem.; (1996) 60, 10, 1720-23 CODEN: BBBIEJ ISSN: 0916-8451
- AU Kobayashi K; Kato M; Miura Y; Kettoku M; Komeda T; Iwamatsu A
- AN 1996-15030 BIOTECHDS
- L61 ANSWER 43 OF 134 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V. on STN DUPLICATE 21
- TI Molecular analysis of the amy gene locus of Thermoanaerobacterium thermosulfurigenes EM1 encoding starch-degrading enzymes and a binding protein-dependent maltose transport system.
- SO Journal of Bacteriology, (1996) 178/4 (1039-1046). ISSN: 0021-9193 CODEN: JOBAAY
- AU Sahm K.; Matuschek M.; Muller H.; Mitchell W.J.; Bahl H.
- AN 96053058 EMBASE
- L61 ANSWER 44 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN DUPLICATE 22
- TI CHARACTERIZATION AND MOLECULAR-CLONING OF THERMOSTABLE ALPHA-AMYLASE FROM STREPTOMYCES SP TO1
- SO BIOTECHNOLOGY LETTERS, (JUL 1996) Vol. 18, No. 7, pp. 809-814. ISSN: 0141-5492.
- AU MELLOULI L; GHORBEL R; KAMMOUN A; MEZGHANI M; BEJAR S (Reprint)
- AN 96:537559 SCISEARCH
- L61 ANSWER 45 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN DUPLICATE 23
- TI Cloning, nucleotide **sequence**, and hyperexpression of **alpha-amylase gene** from an archaeon, Thermococcus profundus
- SO JOURNAL OF FERMENTATION AND BIOENGINEERING, (20 JAN 1996) Vol. 82, No. 5, pp. 432-438.

 Publisher: SOC FERMENTATION BIOENGINEERING, JAPAN, OSAKA UNIV, FACULTY ENGINEERING, 2-1 YAMADAOKA, SUITA, OSAKA 565, JAPAN.

 ISSN: 0922-338X.
- AU Lee J T; Kanai H; Kobayashi T; Akiba T (Reprint); Kudo T
- AN 97:87567 SCISEARCH
- L61 ANSWER 46 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN DUPLICATE 24
- TI CLONING AND EXPRESSION OF THE ALPHA-AMYLASE

 GENE FROM THE HYPERTHERMOPHILIC ARCHAEON PYROCOCCUS SP

KOD1, AND CHARACTERIZATION OF THE ENZYME

JOURNAL OF FERMENTATION AND BIOENGINEERING, (1996) Vol. 82, No. 3, pp. SO 224-232.

ISSN: 0922-338X.

TACHIBANA Y; LECLERE M M; FUJIWARA S; TAKAGI M; IMANAKA T (Reprint) AU

ΑN 96:824456 SCISEARCH

ANSWER 47 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61

TT Cloning and expression of the alpha-amylase

gene from the hyperthermophilic archaeon Pyrococcus sp.

KOD1, and characterization of the enzyme;

thermostable enzyme purification, characterization and gene over-expression from plasmid pET-8c in Escherichia coli

SO J.Ferment.Bioeng.; (1996) 82, 3, 224-32 CODEN: JFBIEX ISSN: 0922-338X

Tachibana Y; Mendez Leclere M; Fujiwara S; Takagi M; *Imanaka T ΑU

AN 1996-15040 BIOTECHDS

ANSWER 48 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61

TΤ Gram negative bacteria of increased competence;

Escherichia coli transformation with vector containing a polynucleotide encoding alpha-amylase gene and competency induction; application in genetic engineering

AU Greener A L

AN1995-08712 BIOTECHDS

PΙ WO 9513388 18 May 1995

ANSWER 49 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61

Gene encoding hyperthermostable alpha-amylase ΤT

of Pyrococcus furiosus;

facilitates use of enzyme in industrial processes when cloned in Escherichia coli

AU Taguchi Y; Nishikawa M; Koyama N; Yamamoto K; Asada K; Kato I

AN 1995-07958 BIOTECHDS

PΙ EP 648843 19 Apr 1995

L61 ANSWER 50 OF 134 MEDLINE on STN DUPLICATE 26

Close evolutionary relatedness among functionally distantly related TImembers of the (alpha/beta)8-barrel glycosyl hydrolases suggested by the similarity of their fifth conserved sequence region.

SO FEBS LETTERS, (1995 Dec 11) 377 (1) 6-8. Journal code: 0155157. ISSN: 0014-5793.

AU Janecek S

AN 96130309 MEDLINE

L61 ANSWER 51 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN

TIPurified Pyrococcus furiosus thermostable alpha-amylase; used for industrial glucopolymer e.g. starch liquefaction at high temperature

ΑN 1994-02942 BIOTECHDS

ΡI EP 577257 5 Jan 1994

ANSWER 52 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61

TINew DNA construct encoding Pyrococcus alpha-amylase;

Pyrococcus woesei and Pyrococcus furiosus recombinant enzyme expression in Bacillus sp. for use in high temperature starch liquefaction or starch saccharification

ΑN 1994-14387 BIOTECHDS

ΡI WO 9419454 1 Sep 1994

ANSWER 53 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61

TINew DNA encoding hyper-thermostable alpha-amylase;

Pyrococcus furiosus recombinant pullulanase, alpha-glucosidase, beta-glucosidase or protease production in Escherichia coli or

Bacillus subtilis

- AN 1994-03550 BIOTECHDS
- PI EP 579360 19 Jan 1994
- L61 ANSWER 54 OF 134 MEDLINE on STN DUPLICATE 28
- TI Cloning of the aapT gene and characterization of its product, alpha-amylase-pullulanase (AapT), from thermophilic and alkaliphilic Bacillus sp. strain XAL601.
- SO APPLIED AND ENVIRONMENTAL MICROBIOLOGY, (1994 Oct) 60 (10) 3764-73. Journal code: 7605801. ISSN: 0099-2240.
- AU Lee S P; Morikawa M; Takagi M; Imanaka T
- AN 95077378 MEDLINE
- L61 ANSWER 55 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
- The gene amyE(TV1) codes for a nonglucogenic alpha-amylase from Thermoactinomyces vulgaris 94-2A in Bacillus subtilis;

gene cloning, characterization and phage EMBL3 expression in Escherichia coli and Bacillus subtilis

- SO Appl.Environ.Microbiol.; (1994) 60, 9, 3381-89 CODEN: AEMIDF
- AU Hofemeister B; Koenig S; Hoang V; Engel J; Mayer G; *Hofemeister J
- AN 1994-12478 BIOTECHDS
- L61 ANSWER 56 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN DUPLICATE 29
- TI SEQUENCE SIMILARITIES AND EVOLUTIONARY RELATIONSHIPS OF MICROBIAL, PLANT AND ANIMAL ALPHA-AMYLASES
- SO EUROPEAN JOURNAL OF BIOCHEMISTRY, (01 SEP 1994) Vol. 224, No. 2, pp. 519-524.
 - ISSN: 0014-2956.
- AU JANECEK S (Reprint)
- AN 94:570329 SCISEARCH
- L61 ANSWER 57 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
- TI Synthesis of branched oligosaccharides from starch by two amylases cloned from Bacillus licheniformis;

maltogenic and thermostable alpha-amylase

gene cloning and expression in Escherichia coli for use in starch liquefaction and saccharification for branched oligosaccharide production

- SO Biosci.Biotechnol.Biochem.; (1994) 58, 2, 416-18 CODEN: BBBIEJ
- AU Kim I C; Yoo S H; Lee S J; Oh B H; Kim J W; *Park K H
- AN 1994-04597 BIOTECHDS
- L61 ANSWER 58 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN DUPLICATE 30
- TI EXPRESSION IN ESCHERICHIA-COLI AND STRUCTURE OF THE GENE ENCODING 4-ALPHA-GLUCANOTRANSFERASE FROM THERMOTOGA-MARITIMA CLASSIFICATION OF MALTODEXTRIN GLYCOSYLTRANSFERASES INTO 2 DISTANTLY RELATED ENZYME SUBFAMILIES
- SO SYSTEMATIC AND APPLIED MICROBIOLOGY, (NOV 1994) Vol. 17, No. 3, pp. 297-305.

ISSN: 0723-2020.

- AU HEINRICH P; HUBER W; LIEBL W (Reprint)
- AN 95:27584 SCISEARCH
- L61 ANSWER 59 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
- TI Recombinant DNA expression in amylase-deficient or protease-deficient Bacillus licheniformis;
 - for production of cyclomaltodextrin-glucanotransferase, pullulanase or glucose-isomerase
- AN 1993-09547 BIOTECHDS
- PI WO 9310248 27 May 1993

L61 ANSWER 60 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN Strain of Bacillus amyloliquefaciens; TIBacillus licheniformis thermostable alpha-amylase gene cloning and expression AN 1994-04770 BIOTECHDS PΤ SU 1788966 15 Jan 1993 1.61 ANSWER 61 OF 134 MEDLINE on STN DUPLICATE 32 Alpha-amylase from the hyperthermophilic archaebacterium Pyrococcus furiosus. Cloning and sequencing of the gene and expression in Escherichia coli. SO JOURNAL OF BIOLOGICAL CHEMISTRY, (1993 Nov 15) 268 (32) 24402-7. Journal code: 2985121R. ISSN: 0021-9258. Laderman K A; Asada K; Uemori T; Mukai H; Taguchi Y; Kato I; Anfinsen C B ΑIJ AN94043280 MEDLINE L61 ANSWER 62 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN ΤI Sequencing of the amylopullulanase (apu) gene of Thermoanaerobacter e thanolicus 39E, and identification of the active site by site-directe d mutagenesis; DNA sequence determination SO J.Biol.Chem.; (1993) 268, 22, 16332-44 CODEN: JBCHA3 ΑU Mathupala S P; Lowe S E; Podkovyrov S M; *Zeikus J G ΑN 1993-11248 BIOTECHDS ANSWER 63 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61 TI Cloning and sequencing of a gene encoding acidophilic amylase from Bacillus acidocaldarius; thermostable alpha-amylase-pullulanase gene DNA sequence SO J.Gen.Microbiol.; (1993) 139, Pt.10, 2399-407 CODEN: JGMIAN Koivula T T; Hemila H; Pakkanen R; Sibakov M; *Palva I AU AN1993-15401 BIOTECHDS L61 ANSWER 64 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN DUPLICATE TIENHANCED PRODUCTION OF ALPHA-AMYLASE IN FED-BATCH CULTURES OF BACILLUS-SUBTILIS TN106[PAT5] SO BIOTECHNOLOGY AND BIOENGINEERING, (20 NOV 1993) Vol. 42, No. 10, pp. 1142-1150. ISSN: 0006-3592. LEE J; PARULEKAR S J (Reprint) ΑU AN93:657450 SCISEARCH L61 ANSWER 65 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN An extremely thermostable alpha-amylase from the ΤI hyperthermophilic archaebacterium Pyrococcus furiosus; thermostable recombinant alpha-amylase production and characterization (conference abstract) SO Protein Eng.; (1993) 6, 8, 1010 CODEN: PRENE9 ΑU Asada K; Mukai H; Uemori T; Taguchi Y; Izu H AN 1994-01082 BIOTECHDS L61 ANSWER 66 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN TIEvaluation of column flotation in the downstream processing of fermen tation products: recovery of a genetically engineered alpha-amylase; Bacillus stearothermophilus recombinant alpha-amylase purification fr om Escherichia coli fermentation broth and periplasmic extract by pha se partitioning and column flotation

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Biotechnol.Prog.; (1993) 9, 4, 411-20

CODEN: BIPRET ΑU Miranda E A; *Berglund K A ΑN 1993-11223 BIOTECHDS L61 ANSWER 67 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN TΙ A neopullulanase-type alpha-amylase gene from Thermoactinomyces vulgaris R-47; gene cloning and DNA sequence determination Biosci.Biotechnol.Biochem.; (1993) 57, 3, 395-401 SO CODEN: BBBIEJ Tonozuka T; Ohtsuka M; Mogi S; Sakai H; Ohta T; *Sakano Y ΑU 1993-05903 BIOTECHDS ΑN L61 ANSWER 68 OF 134 MEDLINE on STN DUPLICATE 34 ΤI Sequence of the Streptomyces thermoviolaceus CUB74 alpha -amylase-encoding gene and its transcription analysis in Streptomyces lividans. GENE, (1993 May 15) 127 (1) 133-7. SO Journal code: 7706761. ISSN: 0378-1119. Bahri S M; Ward J M ΑU 93252270 MEDLINE ΑN L61 ANSWER 69 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN ΤI Construction of inducible secretion vectors and their application for the secretion of foreign extracellular and intracellular proteins in Bacillus subtilis; penicillinase, alpha-amylase, cellulase and beta-galactosidase gene c loning, expression and protein secretion using a plasmid pISA412 or p lasmid pISAts412 vector SO J.Ferment.Bioeng.; (1993) 76, 1, 1-6 CODEN: JFBIEX Imanaka T; Takaqi M; Shima H; Bhatnagar L; Zeikus J G ΑU AN 1993-11370 BIOTECHDS ANSWER 70 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61 Nucleotide sequence of Bacillus stearothermophilus ΤI alpha-amylase gene and its high expression; cloning and DNA sequence SO Ind.Microbiol.; (1993) 23, 2, 1-7 CODEN: GOWEEK Xu Y; Wang X; He C; Wu C; Ren D ΑU AN1994-02920 BIOTECHDS ANSWER 71 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61 ΤI Hybrid alpha-amylase promoter; gene cloning, expression and protein secretion in e.g. Bacillus subtilis using a plasmid pKTH1910, plasmid pKTH1975 or plasmid pKTH1912 vector containing a signal peptide sequence AN1992-06703 BIOTECHDS ΡI WO 9203561 5 Mar 1992 ANSWER 72 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61 Construction of DNA fragment encoding thermostable alpha-amylase; TIBacillus licheniformis gene cloning and expression in Bacillus subtilis, with Bacillus amyloliquefaciens regulatory elements AN 1993-04134 BIOTECHDS SU 1717633 7 Mar 1992 PТ L61 ANSWER 73 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN TICloning of Bacillus licheniformis amylase gene and its application to synthesis of branched oligosaccharides; oligosaccharide preparation and starch liquefaction using panose-forming amylase and recombinant thermostable alpha-amylase expressed in Escherichia coli (conference paper)

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SO Biochem.Eng.2001; (1992) 80-83
AU Park K H; Kim I C; Kim J R; Seo B C; Choi Y D; Lee D S
AN 1993-04106 BIOTECHDS

L61 ANSWER 74 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERI
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ANSWER 74 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN Characterization of the alpha-amylase-encoding

gene from Thermomonospora curvata;

gene cloning and expression in Streptomyces lividans, and DNA sequence SO Gene; (1992) 112, 1, 77-83 CODEN: GENED6

AU Petricek M; *Tichy P; Kuncova M

AN 1992-07671 BIOTECHDS

L61 ANSWER 75 OF 134 HCAPLUS COPYRIGHT 2003 ACS on STN

TI Study of large scale enzyme production method applied for host bacterium, Bacillus brevis

SO Shokuhin Sangyo Senta Gijutsu Kenkyu Hokoku (1992), 18, 43-50 CODEN: SSGHD6; ISSN: 0388-3388

AU Uchida, Kazuhiko; Miyauchi, Akira; Takagi, Hiroaki; Kadowaki, Kiyoshi

AN 1993:21003 HCAPLUS

DN 118:21003

ANSWER 76 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN Gene expression using Gram-positive bacteria; penicillinase, alpha-amylase, neopullulanase, protease, etc. gene cloning and expression in Bacillus stearothermophilus and Bacillus subtilis using a plasmid pTB19 vector (conference paper)

SO Harnessing Biotechnol.21st Century; (1992) 18-22 CODEN: 9999V

AU Imanaka T

AN 1994-02934 BIOTECHDS

L61 ANSWER 77 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN Random point mutation analysis of the signal peptide cleavage area of Bacillus stearothermophilus alpha-amylase;

site-directed mutagenesis and application to improved protein

SO Agric.Biol.Chem.; (1991) 55, 11, 2875-76 CODEN: ABCHA6

AU Yamaguchi K; Ueda M; Kawanishi G; Udaka S

AN 1992-01800 BIOTECHDS

ANSWER 78 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN

Alpha-amylase of Clostridium thermosulfurogenes EM1:
nucleotide sequence of the gene, processing of the enzyme, and comparison to other alpha-amylases;
gene cloning in Escherichia coli; DNA sequence

SO Appl.Environ.Microbiol.; (1991) 57, 5, 1554-59
CODEN: AEMIDF

AU Bahl H; Burchhardt G; Spreinat A; Haeckel K; Wienecke A; Schmidt B AN 1991-08308 BIOTECHDS

L61 ANSWER 79 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN Cloning and expression of an amylase gene from Bacillus stearothermophilus:

thermostable alpha-amylase expression in Bacillus subtilis and Bacillus licheniformis (conference paper)

SO Res.Microbiol.; (1991) 142, 7-8, 793-96 CODEN: RMCREW

AU Diderichsen B; Poulsen G B; Jorgensen P L

AN 1992-01709 BIOTECHDS

L61 ANSWER 80 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN

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Production of thermophilic alpha-amylase using immobilized
        transformed Escherichia coli by addition of glycine;
           immobilization for Bacillus stearothermophilus recombinant
           thermostable alpha-amylase production and plasmid stability
       J.Ferment.Bioeng.; (1991) 71, 6, 397-402
 SO
       CODEN: JFBIEX
 ΑU
       Ariga O; Andoh Y; Fujishita Y; Watari T; Sano Y
       1991-10763 BIOTECHDS
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       ANSWER 81 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
 L61
       Cloning of a chromosomal alpha-amylase gene
 TI
       from Bacillus stearothermophilus;
          amyS gene localization in chromosome: expression in Escherichia coli
 SO
       FEMS Microbiol.Lett.; (1991) 77, 2-3, 271-76
       CODEN: FMLED7
       Jorgensen P L; Poulsen G B; *Diderichsen B
 AII
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       1991-04140 BIOTECHDS
       ANSWER 82 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
 L61
       Effect of cultivation temperatures on {\bf thermophilic} and
 ΤI
       mesophilic enzyme gene expression in Bacillus subtilis;
          Bacillus stearothermophilus thermostable neutral protease and Bacillus
          subtilis alpha-amylase gene cloning and
          expression in B. subtilis
       J.Ferment.Bioeng.; (1991) 72, 3, 193-97
       CODEN: JFBIEX
 ΑU
       Kubo M; Imanaka T
 ΑN
       1991-15032 BIOTECHDS
     ANSWER 83 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN
      EFFECT OF CULTIVATION TEMPERATURES ON THERMOPHILIC AND
TΤ
      MESOPHILIC ENZYME GENE-EXPRESSION IN BACILLUS-SUBTILIS
     JOURNAL OF FERMENTATION AND BIOENGINEERING, (1991) Vol. 72, No. 3, pp.
SO
      193-197.
AU
     KUBO M (Reprint); IMANAKA T
AN
     91:549642 SCISEARCH
      ANSWER 84 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
L61
      Isolation of the pullulanase gene from Clostridium thermosulfurogenes
TΙ
       (DSM 3896) and its expression in Escherichia coli;
          thermostable enzyme gene cloning using plasmid pCT3
SO
      Curr.Microbiol.; (1991) 22, 2, 91-95
      CODEN: CUMIDD
ΑIJ
      Burchhardt G; Wienecke A; *Bahl H
AN
      1991-07140 BIOTECHDS
L61 ANSWER 85 OF 134 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN
     NUCLEOTIDE-SEQUENCE OF 2 CLOSTRIDIUM-THERMOSULFUROGENES EM1 GENES
ΤI
     HOMOLOGOUS TO ESCHERICHIA-COLI GENES ENCODING INTEGRAL MEMBRANE-COMPONENTS
     OF BINDING PROTEIN-DEPENDENT TRANSPORT-SYSTEMS
     FEMS MICROBIOLOGY LETTERS, (1991) Vol. 81, No. 1, pp. 83-88.
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ΑN
     91:365857 SCISEARCH
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L61
      Conjugal transfer of recombinant transposon Tn916 from Escherichia coli
ΤI
      to Bacillus stearothermophilus;
           alpha-amylase gene cloning in vector
         plasmid pAM120A; transposition; conjugation
SO
      Plasmid; (1991) 26, 1, 67-73
      CODEN: PLSMDX
ΑIJ
      Natarajan M R; Oriel P
AN
      1991-13224 BIOTECHDS
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ANSWER 87 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61 TΤ New thermostable enzyme with both alpha-amylase and pullulanase activities; prepared by expressing Clostridium thermohydrosulfuricum DNA in host cells, useful in starch hydrolysis, etc.; DNA sequence AN 1991-02950 BIOTECHDS PΙ EP 402092 12 Dec 1990 ANSWER 88 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN 1.61 ΤI Maltose and maltitol preparation method; starch saccharification using beta-amylase, pullulanase, isoamylase and recombinant alpha-amylase; Bacillus stearothermophilus gene cloning in Bacillus subtilis 1990-07073 BIOTECHDS AN PΙ JP 02042997 13 Feb 1990 ANSWER 89 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61 Expression unit from the replication region of the streptococcal plasmid ΤI pSM19035; Streptococcus pyogenes expression DNA sequence EU19035 isolation and characterization; plasmid pCB22 vector construction; alphaamylase gene cloning in Bacillus subtilis SO Mol.Biol.(Moscow); (1990) 24, 4, 993-1000 CODEN: MOBIBO ΑU Sorokin A V; Khazak V E ΑN 1991-01231 BIOTECHDS L61 ANSWER 90 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN Regulation of thermostable alpha-amylase of Streptomyces thermoviolaceus TICUB74: maltotriose is the smallest inducer; effect of various sugars and C-sources on enzyme induction and repression in the parent strain and of the recombinant enzyme in Streptomyces lividans SO Biochimie; (1990) 72, 12, 893-95 CODEN: BICMBE ΑU Bahri S M; Ward M ΑN 1991-04134 BIOTECHDS ANSWER 91 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61 Cloning and expression of an alpha-amylase TIgene from Streptomyces thermoviolaceus CUB74 in Escherichia coli JM107 and S. lividans TK24; cloning in Streptomyces lividans and protein secretion J.Gen.Microbiol.; (1990) 136, Pt.5, 811-18 SO CODEN: JGMIAN Bahri S M; Ward J M ΑU ΑN 1990-10275 BIOTECHDS ANSWER 92 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN L61 ΤI Engineering of amylase and endoglucanase activity in Lactobacillus plantarum; Bacillus stearothermophilus alpha-amylase and Clostridium thermocellum cellulase gene cloning; application to silage starter culture strain improvement (conference abstract) Food Biotechnol.; (1990) 4, 1, 554 SO CODEN: FBIOEE ΑU Trees S; Jacques M; Henk J; Frank M ΑN 1992-01552 BIOTECHDS 1.61 ANSWER 93 OF 134 LIFESCI COPYRIGHT 2003 CSA on STN DUPLICATE 39 Characterization of a thermostable Bacillus stearothermophilus TI-amylase. SO BIOTECHNOL. APPL. BIOCHEM., (1990) vol. 12, no. 4, pp. 427-435.

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Vihinen, M.; Maentsaelae, P.

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- TI Site-directed mutagenesis of a thermostable alpha-amylase from Bacillus stearothermophilus: putative role of three conserved residues; gene cloning and expression in Escherichia coli; enzyme engineering
- SO J.Biochem.; (1990) 107, 2, 267-72 CODEN: JOBIAO
- AU Vihinen M; Ollikka P; Niskanen J; Meyer P; Suominen I; Karp M
- AN 1990-04728 BIOTECHDS
- L61 ANSWER 95 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
- TI Presence of the bacterial hemoglobin **gene** improves **alpha-amylase** production of a recombinant Escherichia coli strain;

Bacillus stearothermophilus recombinant enzyme preparation; Vitreoscilla sp. hemoglobin gene cloning, vector plasmid pMK79 construction, expression; aerobic growth in microaerophilic conditions

SO Plasmid; (1990) 24, 3, 190-94

CODEN: PLSMDX

- AU Khosravi M; Webster D A; *Stark B C
- AN 1991-02943 BIOTECHDS
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- TI Potential use of Bacillus brevis for enzyme production. ENZYME ENGINEERING 10.
- SO ANN. N.Y. ACAD. SCI., (1990) pp. 582-583.

 Meeting Info.: Tenth International Enzyme Engineering Conference.

 Kashikojima (Japan). 24-29 Sep 1989.

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- AU Udada, S.; Okada, H. [editor]; Tanaka, A. [editor]; Blanch, H.W. [editor]
- AN 90:102234 LIFESCI
- L61 ANSWER 97 OF 134 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT/ISI on STN
- TI Genetics of Streptococcus **thermophilus**: a review; starter culture genetic engineering and strain improvement (conference paper)
- SO J.Dairy Sci.; (1989) 72, 12, 3444-54 CODEN: JDSCAE
- AU Mercenier A; Lemoine Y
- AN 1990-05150 BIOTECHDS
- L61 ANSWER 98 OF 134 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V. on STN DUPLICATE 40
- TI Cloning of a thermostable .alpha.-amylase gene from Thermomonospora curvata and its expression in Streptomyces lividans.
- SO Journal of General Microbiology, (1989) 135/12 (3303-3309). ISSN: 0022-1287 CODEN: JGMIAN
- AU Petricek M.; Stajner K.; Tichy P.
- AN 90034655 EMBASE
- L61 ANSWER 99 OF 134 MEDLINE on STN DUPLICATE 41
- TI Expression of the insecticidal protein gene from Bacillus thuringiensis subsp. aizawai in Bacillus subtilis and in the **thermophile**Bacillus stearothermophilus by using the alpha-amylase promoter of the **thermophile**.
- SO APPLIED AND ENVIRONMENTAL MICROBIOLOGY, (1989 Dec) 55 (12) 3208-13. Journal code: 7605801. ISSN: 0099-2240.
- AU Nakamura K; Imanaka T
- AN 90146333 MEDLINE
- L61 ANSWER 100 OF 134 HCAPLUS COPYRIGHT 2003 ACS on STN
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 2	L2	5392	hyperthermophil\$ or thermophil\$		2003/09/19 14:54
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ABSTRACT:

The invention provides polynucleotides, preferably synthetic polynucleotides, which encode processing enzymes that are optimized for expression in plants. The polynucleotides encode mesophilic, thermophilic, or hyperthermophilic processing enzymes, which are activated under suitable activating conditions to act upon the desired substrate. Also provided are "self-processing" transgenic plants, and plant parts, e.g., grain, which express one or more of these enzymes and have an altered composition that facilitates plant and grain processing. Methods for making and using these plants, e.g., to produce food products having improved taste and to produce fermentable substrates for the production of ethanol and fermented beverages are also provided.

RELATED APPLICATIONS

[0001] This application claims priority to Application Serial No. 60/315,281, filed Aug. 27, 2001, which is herein incorporated by reference.

----- KWIC -----

Summary of Invention Paragraph - BSTX (22):

[0020] Moreover, the present invention encompasses a plant stably transformed with the vectors of the present invention. A plant stably transformed with a vector comprising an anjuna amino acid sequence of any of SEQ ID NO: 1, 10, 13, 14, 15, 16, 33, or 35, or encoded by a polynucleotide comprising any of SEQ ID NO: 2 or 9 is provided. Preferably, the alpha -amylase is hyperthermophilic.

Summary of Invention Paragraph - BSTX (36):

[0034] In a most preferred embodiment, a method of producing hypersweet corn comprising treating transformed corn or a part thereof, the genome of which is augmented with and expresses in the endosperm an expression cassette encoding an .alpha.-amylase, under conditions which activate the at least one enzyme so as to convert polysaccharides in the corn into sugar, yielding hypersweet corn is described. Preferably, the enzyme is hyperthermophilic and the hyperthermophilic .alpha -amylase comprises the amino acid sequence of any of SEQ ID NO: 10, 13, 14, 15, 16, 33, or 35, or an enzymatically active fragment thereof having .alpha.-amylase activity.

Summary of Invention Paragraph - BSTX (38):

[0036] In another aspect of the invention, a method of preparing hydrolyzed starch product comprising treating a plant part comprising starch granules and at least one starch processing enzyme under conditions which activate the at least one enzyme thereby processing the starch granules to form an aqueous solution comprising a hydrolyzed starch product, wherein the plant part is obtained from a transformed plant, the genome of which is augmented with an expression cassette encoding at least one .alpha.-amylase; and collecting the aqueous solution comprising hydrolyzed starch product is described. Preferably, the .alpha.-amylase is hyperthermophilic and more preferably, the hyperthermophilic .alpha.-amylase comprises the amino acid sequence of any of SEQ ID NO: 1, 10, 13, 14, 15, 16, 33, or 35, or an active fragment thereof having .alpha.-amylase activity. Preferably, the expression cassette comprises a polynucleotide selected from any of SEQ ID NO: 2, 9, 46, or 52, a complement thereof, or a polynucleotide that hybridizes to any of SEQ ID NO: 2, 9, 46, or 52 under low stringency hybridization conditions and encodes a polypeptide having .alpha.-amylase activity. Moreover, the invention further provides for the genome of the transformed plant further comprising a polynucleotide encoding a non-thermophilic starch-processing enzyme. Alternatively, the plant part may be treated with a non-hyperthermophilic starch-processing enzyme.

Detail Description Paragraph - DETX (33):

[0108] In another embodiment of the invention, the polynucleotide encodes a hyperthermophilic processing enzyme that is operably linked to a chloroplast (amyloplast) transit peptide (CTP) and a starch binding domain, e.g., from the waxy gene. An exemplary polynucleotide in this embodiment encodes SEQ ID NO:10 .alpha.-amylase linked to the starch binding domain from waxy). Other exemplary polynucleotides encode a hyperthermophilic processing enzyme linked to a signal sequence that targets the enzyme to the endoplasmic reticulum and

secretion to the apoplast (exemplified by a polynucleotide encoding SEQ ID NO:13, 27, or 30, which comprises the N-terminal sequence from maize .gamma.-zein operably linked to .alpha.-amylase, .alpha.-glucosidase, glucose isomerase, respectively), a hyperthermophilic processing enzyme linked to a signal sequence which retains the enzyme in the endoplasmic reticulum (exemplified by a polynucleotide encoding SEQ ID NO:14, 26, 28, 29, 33, 34, 35, or 36, which comprises the N-terminal sequence from maize .gamma.-zein operably linked to the <u>hyperthermophilic</u> enzyme, which is operably linked to SEKDEL, wherein the enzyme is .alpha.-amylase, malA .alpha.-glucosidase, T. maritima glucose isomerase, T. neapolitana glucose isomerase), a hyperthermophilic processing enzyme linked to an N-terminal sequence that targets the enzyme to the amyloplast (exemplified by a polynucleotide encoding SEQ ID NO:15, which comprises the N-terminal amyloplast targeting sequence from waxy operably linked to .alpha.-amylase), a hyperthermophilic fusion polypeptide which targets the enzyme to starch granules (exemplified by a polynucleotide encoding SEQ ID NO:16, which comprises the N-terminal amyloplast targeting sequence from waxy operably linked to an .alpha.-amylase/waxy fusion polypeptide comprising the waxy starch binding domain), a hyperthermophilic processing enzyme linked to an ER retention signal (exemplified by a polynucleotide encoding SEQ ID NO:38 and 39). Moreover, a <u>hyperthermophilic</u> processing enzyme may be linked to a raw-starch binding site having the amino acid sequence (SEQ ID NO:53), wherein the polynucleotide encoding the processing enzyme is linked to the maize-optimized nuleic acid sequence (SEQ ID NO:54) encoding this binding site.

Detail Description Paragraph - DETX (40):

[0115] A signal sequence such as the maize .gamma.-zein N-terminal signal sequence for targeting to the endoplasmic reticulum and secretion into the apoplast may be operably linked to a polynucleotide encoding a hyperthermophilic processing enzyme in accordance with the present invention (Torrent et al., 1997). For example, SEQ ID NOs: 13, 27, and 30 provides for a polynucleotide encoding a hyperthermophilic enzyme operably linked to the N-terminal sequence from maize .gamma.-zein protein. Another signal sequence is the amino acid sequence SEKDEL for retaining polypeptides in the endoplasmic reticulum (Munro and Pelham, 1987). For example, a polynucleotide encoding SEQ ID NOS:14, 26, 28, 29, 33, 34, 35, or 36, which comprises the N-terminal sequence from maize .gamma -zein operably linked to a processing enzyme which is operably linked to SEKDEL. A polypeptide may also be targeted to the amyloplast by fusion to the waxy amyloplast targeting peptide (Klosgen et al., 1986) or to a starch granule. For example, the polynucleotide encoding a hyperthermophilic processing enzyme may be operably linked to a chloroplast (amyloplast) transit peptide (CTP) and a starch binding domain, e.g., from the waxy gene. SEQ ID NO:10 exemplifies .alpha.-amylase linked to the starch binding domain from waxy. SEQ ID NO:15 exemplifies the N-terminal sequence amyloplast targeting sequence from waxy operably linked to .alpha.-amylase. Moreover, the polynucleotide encoding the processing enzyme may be fused to target starch granules using the waxy starch binding domain. For example, SEQ ID NO:16 exemplifies a fusion polypeptide comprising the N-terminal amyloplast targeting sequence from waxy operably linked to an .alpha.-amylase/waxy fusion polypeptide comprising the waxy starch binding domain.

Detail Description Paragraph - DETX (118):

[0191] The invention provides a method to produce dextrins and altered starches from a plant, or a product from a plant, that has been transformed with a processing enzyme which hydrolyses certain covalent bonds of a polysaccharide to form a polysaccharide derivative. In one embodiment, a plant, or a product of the plant such as a fruit or grain, or flour made from the grain that expresses the enzyme is placed under conditions sufficient to activate the enzyme and convert polysaccharides contained within the plant into polysaccharides of reduced molecular weight. Preferably, the enzyme is fused to a signal sequence that targets the enzyme to a starch granule, an amyloplast, the apoplast or to the endoplasmic reticulum as disclosed herein. The dextrin or derivative starch produced may then be isolated or recovered from the plant or the product of the plant. In another embodiment, a processing enzyme able to convert polysaccharides into dextrins or altered starches is placed under the control of an inducible promoter according to methods known in the art and disclosed herein. The plant is grown to a desired stage and the promoter is induced causing expression of the enzyme and conversion of the polysaccharides, within the plant or product of the plant, to dextrins or altered starches. Preferably the enzyme is .alpha.-amylase. pullulanase, iso or neo-pullulanase and is operably linked to a signal sequence that targets the enzyme to a starch granule, an amyloplast, the apoplast or to the endoplasmic reticulum. In one embodiment, the enzyme is targeted to the apoplast or to the endoreticulum. In yet another embodiment, a transformed plant is produced that expresses an enzyme able to convert starch into dextrins or altered starches. The enzyme is fused to a signal sequence that targets the enzyme to a starch granule within the plant. Starch is then isolated from the transformed plant that contains the enzyme expressed by the transformed plant. The enzyme contained in the isolated starch may then be activated under conditions sufficient for activation to convert the starch into dextrins or altered starches. Examples of hyperthermophilic enzymes, for example, able to convert starch to hydrolyzed starch products are provided herein. The methods may be used with any plant which produces a polysaccharide and that can express an enzyme able to convert a polysaccharide into sugar.

Detail Description Paragraph - DETX (123):

[0196] The invention also provides for the production of improved corn varieties (and varieties of other crops) that have normal levels of starch accumulation, and accumulate sufficient levels of amylolytic enzyme(s) in their endosperm, or starch accumulating organ, such that upon activation of the enzyme contained therein, such as by boiling or heating the plant or a part thereof in the case of a hyperthermophilic enzyme, the enzyme(s) is activated and facilitates the rapid conversion of the starch into simple sugars. These simple sugars (primarily glucose) will provide sweetness to the treated corn. The resulting corn plant is an improved variety for dual use as a grain producing hybrid and as sweet corn. Thus, the invention provides a method to produce hyper-sweet corn, comprising treating transformed corn or a part thereof, the genome of which is augmented with and expresses in endosperm an expression cassette comprising a promoter operably linked to a first polynucleotide encoding at least one amylolytic enzyme, conditions which activate the at least one enzyme so as to convert polysaccharides in the corn into sugar, yielding hypersweet corn. The promoter may be a constitutive promoter, a seed-specific promoter, or an endosperm-specific promoter which is

linked to a polynucleotide sequence which encodes a processing enzyme such as .alpha.-amylase, e.g., one comprising SEQ ID NO:13, 14, or 16. Preferably, the enzyme is <u>hyperthermophilic</u>. In one embodiment, the expression cassette further comprises a second polynucleotide which encodes a signal sequence operably linked to the enzyme encoded by the first polynucleotide. Exemplary signal sequences in this embodiment of the invention direct the enzyme to apoplast, the endoplasmic reticulum, a starch granule, or to an amyloplast. The corn plant is grown such that the ears with kernels are formed and then the promoter is induced to cause the enzyme to be expressed and convert polysaccharide contained within the plant into sugar.

Detail Description Paragraph - DETX (162):

[0232] The 797GL3 _alpha.-amylase, having the amino acid sequence SEQ ID NO:1, was selected for its hyperthermophilic activity. This enzyme's nucleic acid sequence was deduced and maize-optimized as represented in SEQ ID NO:2. Similarly, the 6gp3 pullulanase was selected having the amino acid sequence set forth in SEQ ID NO:3. The nucleic acid sequence for the 6gp3 pullulanase was deduced and maize-optimized as represented in SEQ ID NO:4.

Claims Text - CLTX (136):

135. The method of claim 134, wherein the hyperthermophilic .alpha.-amylase comprises the amino acid sequence of any of SEQ ID NO:10, 13, 14, 15, 16, 33, or 35, or an enzymatically active fragment thereof having .alpha.-amylase activity.

Claims Text - CLTX (153):

152. The method of claim 151, wherein the hyperthermophilic .alpha.-amylase comprises the amino acid sequence of any of SEQ ID NO: 1, 10, 13, 14, 15, 16, 33, or 35, or an active fragment thereof having .alpha.-amylase activity.

PGPUB-DOCUMENT-NUMBER: 20030125534

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030125534 A1

Enzymes having alpha amylase activity and methods of TITLE:

use thereof

July 3, 2003 PUBLICATION-DATE:

INVENTOR-INFORMATION:

COUNTRY RULE-47 STATE CITY NAME

US CA San Diego Callen, Walter US CA San Diego Richardson, Toby US CA San Diego Frey, Gerhard US Rancho Santa Fe CA Short, Jay M. US CA Carlsbad Mathur, Eric J. US CA San Diego Gray, Kevin A. US CA San Diego Kerovuo, Janne E. US CA San Diego Slupska, Malgorzata

10/081872 APPL-NO:

DATE FILED: February 21, 2002

RELATED-US-APPL-DATA:

non-provisional-of-provisional 60270495 20010221 US

non-provisional-of-provisional 60270496 20010221 US

non-provisional-of-provisional 60291122 20010514 US

US-CL-CURRENT: 536/23.1

ABSTRACT:

The invention relates to alpha amylases and to polynucleotides encoding the alpha amylases. In addition methods of designing new alpha amylases and methods of use thereof are also provided. The alpha amylases have increased activity and stability at acidic, neutral and alkaline pH and increased temperature.

RELATED APPLICATION DATA

[0001] This application claims priority of U.S. Provisional Application No. 60/270,495, filed Feb. 21, 2001, now pending; U.S. Provisional Application No. 60/270,496, filed Feb. 21, 2001, now pending, and U.S. Provisional Application No. 60/291,122, filed May 14, 2001, now pending, all of which are herein incorporated by reference in their entirety.

KWIC

Detail Description Paragraph - DETX (516):

[0547] An Initial bioinformatic analysis was made with the known hyperthermophillic .alpha.-amylase sequences. FIG. 14a shows an alignment of the sequences some of which have been deposited at the NCBI database. This analysis revealed the potential for designing degenerate primers to PCR the entire gene minus its signal sequence (see FIG. 14a), yielding potentially novel full-length alpha amylases from a library.

PGPUB-DOCUMENT-NUMBER: 20020106779

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020106779 A1

TITLE:

Thermostable peptidase

PUBLICATION-DATE:

August 8, 2002

INVENTOR-INFORMATION:

CITY NAME

STATE COUNTRY RULE-47 US

Cheng, Timothy C.

CA Pasadena Pasadena

Ramakrishnan, Vij

US CA

Chan, Sunney I.

Pasadena

US CA

APPL-NO:

09/969125

DATE FILED: September 24, 2001

RELATED-US-APPL-DATA:

child 09969125 A1 20010924

parent division-of 09333768 19990615 US GRANTED

parent-patent 6294367 US

non-provisional-of-provisional 60089398 19980615 US

US-CL-CURRENT: 435/226, 435/252.3 , 435/320.1 , 435/69.1 , 536/23.2

ABSTRACT:

Thermostable peptidase enzyme derived from archaeon from the genus Pyrococcus is disclosed. The enzyme is produced from native or recombinant host cells and can be utilized in the biotechnology industry as a useful enzyme in sequencing reactions.

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application claims priority from Provisional Application Serial No. 60/089,398, filed Jun. 15, 1998, which is incorporated herein by reference in its entirety and to which application a priority claim is made under 35 U.S.C. .sctn.119(e).

 KWIC	
 114410	

Summary of Invention Paragraph - BSTX (5):

[0005] Most of the proteins isolated from these hyperthermophiles exhibit a temperature optimum of at least 80-100.degree. C. or above (Adams et al., Bio/Technology 13, 662-668 (1995); Adams et al., Trends Biotechnol 16, 329-332 (1998)). Accordingly, there is much interest in exploiting these proteins for biotechnological applications, as they are able to perform biochemical reactions under harsh conditions, such as in the presence of high-temperatures, organic solvents, and denaturants (Adams et al., supra.) P. furiosus has been the source of many of these biotechnologically important proteins, including DNA polymerase (Lundberg et al., Gene 108, 1-6 (1991)), alpha.-amylase (Laderman et al., J. Biol. Chem. 268, 24394-24401 (1993)), and proteases (Voorhorst et al., J. Biol. Chem. 271, 20426-20431 (1996); Harwood et al., J. Bacteriol. 179, 3613-3618 (1997)).

US-PAT-NO:

6426211

DOCUMENT-IDENTIFIER: US 6426211 B1

TITLE:

Xylanase derived from a Bacillus species, expression vectors for such xylanase and other proteins, host

organisms therefor and use thereof

DATE-ISSUED:

July 30, 2002

INVENTOR-INFORMATION:

ZIP CODE COUNTRY STATE CITY NAME N/A BE N/A Linkebeek de Buyl; Eric BE N/A N/A Brussels Lahaye; Andree BE N/A N/A Brussels Ledoux; Pierre N/A BE N/A Rixensart Amory; Antoine BE N/A N/A Detroz; Rene Ohain N/A BE N/A Grez-Doiceau Andre; Christophe DE N/A N/A Burgdorf Vetter; Roman

APPL-NO:

09/073055

DATE FILED: May 5, 1998

PARENT-CASE:

This application is a divisional of 08/275,526 Jul. 15, 1994 now U.S. Pat. No. 6,180,382, Jan. 30, 2001.

FOREIGN-APPL-PRIORITY-DATA:

COUNTRY

APPL-NO

APPL-DATE

GB

9314780

July 15, 1993

US-CL-CURRENT: 435/278, 435/183, 435/200, 435/252.3, 435/262, 435/267 , 435/274 , 435/277 , 435/320.1 , 435/69.1

ABSTRACT:

A purified xylanase derived from B. Pumilus PRL B12 is disclosed. This xylanase is efficient for use in the biobleaching of wood pulp, permitting a strong reduction in the quantity of chlorine used and AOX compounds produced in classical and ECF wood pulp bleaching sequences as well as the quantity of ozone used in TCF sequences. The gene coding for the xylanase was isolated and purified and used to construct an expression vector therefor. A recombinant host strain of B. licheniformis is also disclosed which is efficient for expressing heterologous enzymes, including the xylanase when transformed by the expression vector.

16 Claims, 14 Drawing figures

Exemplary Claim Number:	1
Number of Drawing Sheets:	14
KWIC	

Detailed Description Text - DETX (68):

Other expression vectors provided herein include the nucleotide sequence that codes for the pullulanase of Bacillus deramificans T 89.117D (pUBDEBRA1), or the nucleotide sequence that codes for the .alpha.-amylase of B. licheniformis ATCC 9789 (pL7TAKA), or the nucleotide sequence that codes for the alkaline protease of B. licheniformis SE2 (pLI1), or the subtilisin (alkaline protease) of Bacillus subtilis 168 (pKAC1 and pL7SBT). The expression hosts of the present invention are strains of the genus Bacillus which are compatible with the expression vector for the protein desired to be expressed thereby. Preferably, these strains are aerobic. It is further preferred that these strains not be thermophilic. Such strains include B. subtilis, B. pumilus, and B. licheniformis, B. alkalophilus, B. lentus and B. amyloliquefaciens. Preferably, the alkaline protease gene(s) thereof has (have) been deleted from these expression hosts.

6423523

DOCUMENT-IDENTIFIER: US 6423523 B1

TITLE:

Xylanase derived from a bacillus species, expression vectors for such xylanase and other proteins, host organisms therefor and use thereof

DATE-ISSUED:

July 23, 2002

INVENTOR-INFORMATION:

ZIP CODE COUNTRY STATE CITY NAME BE N/A N/A Linkebeek de Buyl; Eric BE N/A N/A Brussels Lahave: Andree BE N/A N/A Brussels Ledoux: Pierre BE N/A N/A Rixensart Amory; Antoine N/A BE Ohain N/A Detroz; Rene BE N/A N/A Grez-Doiceau Andre; Christophe DE N/A N/A Burgdorf Vetter; Roman

APPL-NO:

09/076677

DATE FILED: May 12, 1998

PARENT-CASE:

This application is a divisional of Ser. No. 08/275,526 filed Jul. 15, 1994 now U.S. Pat. No. 6,180,382, Jan. 30, 2001.

FOREIGN-APPL-PRIORITY-DATA:

COUNTRY

APPL-NO

APPL-DATE

GB

9314780

July 15, 1993

US-CL-CURRENT: 435/200, 435/183, 435/194, 435/252.3, 435/320.1, 435/69.1 , 536/23.2

ABSTRACT:

A purified xylanase derived from B. Pumilus PRL B12 is disclosed. This xylanase is efficient for use in the biobleaching of wood pulp, permitting a strong reduction in the quantity of chlorine used and AOX compounds produced in classical and ECF wood pulp bleaching sequences as well as the quantity of ozone used in TCF sequences. The gene coding for the xylanase was isolated and purified and used to construct an expression vector therefor. A recombinant host strain of B. licheniformis is also disclosed which is efficient for expressing heterologous enzymes, including the xylanase when transformed by the expression vector.

25 Claims, 14 Drawing figures

Exemplary Claim Number: Number of Drawing Sheets:	9
KWIC	

Detailed Description Text - DETX (68):

Other expression vectors provided herein include the nucleotide sequence that codes for the pullulanase of Bacillus deramificans T 89.117D (pUBDEBRA1), or the nucleotide sequence that codes for the .alpha.-amylase of B. licheniformis ATCC 9789 (pL7TAKA), or the nucleotide sequence that codes for the alkaline protease of B. licheniformis SE2 (pLI1I), or the subtilisin (alkaline protease) of Bacillus subtilis 168 (pKAC1 and pL7SBT). The expression hosts of the present invention are strains of the genus Bacillus which are compatible with the expression vector for the protein desired to be expressed thereby. Preferably, these strains are aerobic. It is further preferred that these strains not be thermophilic. Such strains include B. subtilis, B. pumilus, and B. licheniformis, B. alkalophilus, B. lentus and B. amyloliquefaciens. Preferably, the alkaline protease gene(s) thereof has (have) been deleted from these expression hosts.

6329187

DOCUMENT-IDENTIFIER: US 6329187 B1

TITLE:

Endoglucanases

DATE-ISSUED:

December 11, 2001

INVENTOR-INFORMATION:

NAME

STATE

ZIP CODE COUNTRY

Lam; David E.

Harbor City

CA

N/A N/A

Mathur; Eric J.

Carlsbad

CA

N/A N/A

APPL-NO:

09/430669

DATE FILED: October 28, 1999

PARENT-CASE:

This application is a divisional of application Ser. No. 09/066,544 filed on Apr. 24, 1998, now U.S. Pat. No. 6,001,984, which is a continuation application of U.S. application Ser. No. 08/651,572, filed May 22, 1996, now issued as U.S. Pat. No. 5,789,228, the entire contents of which are hereby incorporated by reference herein.

US-CL-CURRENT: 435/209, 435/183, 435/200, 435/220, 536/23.2

ABSTRACT:

A purified thermostable enzyme is derived from the archael bacterium AEPII1a. The enzyme has a molecular weight of about 60.9 kilodaltons and has cellulase activity. The enzyme can be produced from native or recombinant host cells and can be used to aid in the digestion of cellulose where desired.

8 Claims, 2 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 2

----- KWIC -----

Other Reference Publication - OREF (24):

Tachibana Yoshihisa et al., "Cloning and Expression of the Alpha-Amylase Gene from the Hyperthermophilic Archaeon Pyrococcus SP. KOD1, and Characterization of the Enzyme", Journal of Fermentation and Bioengineering, vol. 82, No. 3, 1996 (pp. 224-232).

6300115

DOCUMENT-IDENTIFIER: US 6300115 B1

TITLE:

Pullulanase expression constructs containing .alpha.-amylase promoter and leader sequences

DATE-ISSUED:

October 9, 2001

INVENTOR-INFORMATION:

ZIP CODE COUNTRY NAME STATE

N/A N/A Teague; W. Martin Rockford IL Brumm; Phillip J. Rockford IL N/A N/A Northfield Allen; Larry N. IL N/A N/A Brikun; Igor A. Forest Park IL N/A N/A

APPL-NO:

09/313677

DATE FILED: May 18, 1999

PARENT-CASE:

Priority is claimed to provisional patent application Ser. No. 60/122,065, filed May 18, 1998, and incorporated herein by reference.

US-CL-CURRENT: 435/210, 435/252.31, 435/254.11, 435/320.1, 435/325 , 435/419 , 536/23.1 , 536/23.2 , 536/23.4 , 536/24.1

ABSTRACT:

Disclosed herein are DNA expression constructs containing an .alpha.-amylase promoter sequence derived from Bacillus stearothermophilus, an .alpha.-amylase leader sequence derived from Bacillus stearothermophilus, and a DNA sequence encoding a pullulanase derived from Bacillus naganoensis. Microbial hosts transformed to contain the expression constructs secret function pullulanases. Also disclosed is a process for making recombinant pullulanases utilizing the expression constructs and a recombinant pullulanase which can be produced in Bacillus subtilis.

25 Claims, 6 Drawing figures Exemplary Claim Number: Number of Drawing Sheets: 6 ----- KWIC -----

Other Reference Publication - OREF (23):

Gray et al., Structural <u>Genes Encoding the Thermophilic .alpha.-Amylases</u> of Bacillus stearothermophilus and Bacillus licheniformis, Journal of Bacteriology (May 1986), 166:635-643.

6294367

DOCUMENT-IDENTIFIER: US 6294367 B1

TITLE:

Thermostable peptidase

DATE-ISSUED:

September 25, 2001

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE COUNTRY

Cheng; Timothy C.

Pasadena

N/A N/A

Ramakrishnan; Vij

Pasadena

N/A CA

CA

N/A

Chan; Sunney I.

Pasadena

CA N/A N/A

APPL-NO:

09/333768

DATE FILED: June 15, 1999

PARENT-CASE:

CROSS REFERENCE TO RELATED APPLICATIONS

This application claims priority from Provisional Application Ser. No. 60/089,398, filed Jun. 15, 1998, which is incorporated herein by reference in its entirety and to which application a priority claim is made under 35 U.S.C. .sctn.119(e).

US-CL-CURRENT: 435/212, 435/252.3, 435/320.1, 435/325, 435/455, 435/6 , 435/69.1 , 536/23.2

ABSTRACT:

Thermostable peptidase enzyme derived from archaeon from the genus Pyrococcus is disclosed. The enzyme is produced from native or recombinant host cells and can be utilized in the biotechnology industry as a useful enzyme in sequencing reactions.

19 Claims, 18 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 12

----- KWIC -----

Brief Summary Text - BSTX (5):

Most of the proteins isolated from these **hyperthermophiles** exhibit a temperature optimum of at least 80-100.degree. C. or above (Adams et al., Bio/Technology 13, 662-668 (1995); Adams et al., Trends Biotechnol 16, 329-332 (1998)). Accordingly, there is much interest in exploiting these proteins for biotechnological applications, as they are able to perform biochemical reactions under harsh conditions, such as in the presence of high-temperatures, organic solvents, and denaturants (Adams et al., supra.) P. furiosus has been the source of many of these biotechnologically important proteins, including DNA polymerase (Lundberg et al., Gene 108, 1-6 (1991)), .alpha.-amylase (Laderman et al., J. Biol. Chem. 268, 24394-24401 (1993)), and proteases (Voorhorst et al., J. Biol. Chem. 271, 20426-20431 (1996); Harwood et al., J. Bacteriol. 179, 3613-3618 (1997)).

6218164

DOCUMENT-IDENTIFIER: US 6218164 B1

TITLE:

Thermopallium bacteria and enzymes obtainable therefrom

DATE-ISSUED:

April 17, 2001

INVENTOR-INFORMATION:

NAME

STATE ZIP CODE COUNTRY N/A

Jones: Brian E. Leidschendam Herweijer; Margareta A. The Hague Danson; Michael J. Saltford Hough; David W. Bath

N/A N/A N/A GB N/A N/A GB

N/A

N/A

NL

NL

Thompson; Carl R.

Bath

N/A N/A GB

APPL-NO:

09/029937

DATE FILED: June 2, 1998

FOREIGN-APPL-PRIORITY-DATA:

COUNTRY

APPL-NO

APPL-DATE

EΡ

95202477

September 13, 1995

PCT-DATA:

APPL-NO: PCT/EP96/03896 DATE-FILED: September 3, 1996

PUB-NO: WO97/10342 PUB-DATE: Mar 20, 1997 371-DATE: Jun 2, 1998 102(E)-DATE:Jun 2, 1998

435/210, 435/201, 435/277, 435/278, 435/98, 510/300 US-CL-CURRENT:

, 510/305

ABSTRACT:

The present invention provides thermophilic alkaliphilic bacteria designated Thermopallium natronophilum and thermophilic alkaliphilic polypeptides obtainable therefrom. It also provides compositions, particularly detergent compositions comprising the polypeptides.

10 Claims, 0 Drawing figures

Exemplary Claim Number:

----- KWIC -----

Other Reference Publication - OREF (2):

Lee, S.-P., et al., Applied and Environmental Microbiology, vol. 60, Cloning of the aapT gene and characterization of its product, alpha-amylase-pullulanase (Aapt) from thermophilic and alkaliphilic Bacillus sp. strain XAL601, pp. 3764-3773, 1994.*

6187576

DOCUMENT-IDENTIFIER: US 6187576 B1

TITLE:

.alpha.-amylase mutants

DATE-ISSUED:

February 13, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Svendsen; Allan Birker.o slashed.d N/A N/A DK Borchert; Torben Vedel Jyllinge N/A N/A DK

Bisg.ang.rd-Frantzen; Henrik Bagsv.ae butted.rd N/A N/A DK

APPL-NO: 09/ 170670

DATE FILED: October 13, 1998

PARENT-CASE:

CROSS-REFERENCE TO RELATED APPLICATIONS

The application claims priority under 35 U.S.C 119 of Danish application 1172/97 filed Oct. 13, 1997, and of U.S. provisional application 60/063,306 filed Oct. 28, 1997, the contents of which are fully incorporated herein by reference.

FOREIGN-APPL-PRIORITY-DATA:

COUNTRY

APPL-NO

APPL-DATE

DK

1172/97

October 13, 1997

US-CL-CURRENT: 435/202, 435/183, 435/200, 510/226, 510/235, 510/320, 510/392

ABSTRACT:

The invention relates to a variant of a parent Termamyl-like .alpha.-amylase, comprising mutations in two, three, four, five or six regions/positions. The variants have increased thermostability at acidic pH and/or at low Ca.sup.2+ concentrations (relative to the parent). The invention also relates to a DNA construct comprising a DNA sequence encoding an .alpha.-amylase variant of the invention, a recombinant expression vector which carries a DNA construct of the invention, a cell which is transformed with a DNA construct of the invention, the use of an .alpha.-amylase variant of the invention for washing and/or dishwashing, textile desizing, starch liquefaction, a detergent additive comprising an .alpha.-amylase variant of the invention, a manual or automatic dishwashing detergent composition comprising an .alpha.-amylase variant of the invention, a method for generating a variant of a parent Termamyl-like .alpha.-amylase, which variant exhibits increased thermostability at acidic pH and/or at low Ca.sup.2+ concentrations (relative

to the parent).	
22 Claims, 1 Drawing figure	s
Exemplary Claim Number:	1
Number of Drawing Sheets:	3
KWIC	

Other Reference Publication - OREF (1):
Gray G.L. et al. Structural genes encoding the thermophilic alpha-amylases of Bacillus stearothermophilus and B.licheniformis. J.Bacteriol., May 1986, vol. 166(2):635-643.

6180382

DOCUMENT-IDENTIFIER: US 6180382 B1

TITLE:

Xylanase derived from a bacillus species, expression

vectors for such xylanase and other proteins, host

organisms therefor and use thereof

DATE-ISSUED:

January 30, 2001

INVENTOR-INFORMATION:

STATE ZIP CODE COUNTRY NAME CITY

De Buyl; Eric B-1630 Linkebeek N/A N/A BE N/A N/A BE B-1020 Brussels Lahaye: Andree B-1200 Brussels N/A ΒE Ledoux; Pierre N/A N/A N/A BE B-1330 Rixensart Amory: Antoine BE Detroz: Rene B-1328 Ohain N/A N/A Andre; Christophe B-1390 Grez-Doiceau N/A N/A BE DE

Vetter; Roman

N/A

W-31303 Burgdorf

N/A

08/ 275526

DATE FILED: July 15, 1994

APPL-NO:

FOREIGN-APPL-PRIORITY-DATA:

COUNTRY

APPL-NO

APPL-DATE

GB

9314780

July 15, 1993

US-CL-CURRENT: 435/200, 435/183, 435/201, 435/220, 435/222

ABSTRACT:

A purified xylanase derived from B. Pumilus PRL B12 is disclosed. This xylanase is efficient for use in the biobleaching of wood pulp, permitting a strong reduction in the quantity of chlorine used and AOX compounds produced in classical and ECF wood pulp bleaching sequences as well as the quantity of ozone used in TCF sequences. The gene coding for the xylanase was isolated and purified and used to construct an expression vector therefor. A recombinant host strain of B. licheniformis is also disclosed which is efficient for expressing heterologous enzymes, including the xylanase when transformed by the expression vector.

19 Claims, 14 Drawing figur	es
Exemplary Claim Number:	1
Number of Drawing Sheets:	9
KWIC	

Detailed Description Text - DETX (73):

Other expression vectors provided herein include the nucleotide sequence that codes for the pullulanase of Bacillus deramificans T 89.117D (pUBDEBRA1), or the nucleotide sequence that codes for the .alpha.-amylase of B. licheniformis ATCC 9789 (pL7TAKA), or the nucleotide sequence that codes for the alkaline protease of B. licheniformis SE2 (pLI1), or the subtilisin (alkaline protease) of Bacillus subtilis 168 (pKAC1 and pL7SBT). The expression hosts of the present invention are strains of the genus Bacillus which are compatible with the expression vector for the protein desired to be expressed thereby. Preferably, these strains are aerobic. It is further preferred that these strains not be thermophilic. Such strains include B. subtilis, B. pumilus, and B. licheniformis, B. alkalophilus, B. lentus and B. amyloliquefaciens. Preferably, the alkaline protease gene(s) thereof has (have) been deleted from these expression hosts.

6080568

DOCUMENT-IDENTIFIER: US 6080568 A

TITLE:

Mutant .alpha.-amylase comprising modification at residues corresponding to A210, H405 and/or T412 in

Bacillus licheniformis

DATE-ISSUED:

June 27, 2000

INVENTOR-INFORMATION:

NAME

STATE ZIP CODE COUNTRY

Day; Anthony G. Swanson; Barbara A. San Francisco

N/A N/A CA

San Francisco

CA N/A N/A

APPL-NO:

08/914679

DATE FILED: August 19, 1997

US-CL-CURRENT: 435/202, 435/201, 435/203, 435/275, 435/440, 435/832

, 435/836 , 510/320 , 570/226 , 570/235

ABSTRACT:

Novel .alpha.-amylase enzymes are disclosed in which one or more of residues corresponding to A210, H405 and T412 in Bacillus licheniformis are mutated. The disclosed .alpha.-amylase enzymes show altered or improved stability and/or activity profiles.

11 Claims, 9 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 9

----- KWIC -----

Other Reference Publication - OREF (22):

Gray et al., "Structural Genes Encoding the Thermophilic .alpha.-amylases of Bacillus stearothermophilus and Bacillus licheniformis," J Bacteriol (1986) 166:635-643.

6022724

DOCUMENT-IDENTIFIER: US 6022724 A

TITLE:

.alpha.-amylase mutants

DATE-ISSUED:

February 8, 2000

INVENTOR-INFORMATION:

NAME

ZIP CODE COUNTRY STATE

Svendsen: Allan Birkeroed Bisg.ang.rd-Frantzen; Henrik Lyngby N/A N/A DK N/A DK N/A

Borchert; Torben

Copenhagen N

DK N/A N/A

APPL-NO:

08/683838

DATE FILED: July 18, 1996

PARENT-CASE:

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation-in-part of Ser. No. 08/600,908 filed Feb. 13, 1996 which is a 371 of PCT/DK96/00057 filed Feb. 5, 1996, which are incorporated herein by reference.

FOREIGN-APPL-PRIORITY-DATA:

COUNTRY	APPL-NO	APPL-DATE
DK	0128/95	February 3, 1995
DK	1192/95	October 23, 1995
DK	1256/95	November 10, 1995

US-CL-CURRENT: 435/202, 435/203, 510/226, 510/235, 510/320, 510/392

ABSTRACT:

The present invention relates to a method of constructing a variant of a parent Termamyl-like .alpha.-amylase, which variant has .alpha.-amylase activity and at least one altered property as compared to the parent .alpha.-amylase, comprises

- i) analyzing the structure of the parent Termamyl-like .alpha.-amylase to identify at least one amino acid residue or at least one structural part of the Termamyl-like .alpha.-amylase structure, which amino acid residue or structural part is believed to be of relevance for altering the property of the parent Termamyl-like .alpha.-amylase (as evaluated on the basis of structural or functional considerations),
- ii) constructing a Termamyl-like .alpha.-amylase variant, which as compared to the parent Termamyl-like .alpha.-amylase, has been modified in the amino

acid residue or structural part identified in i) so as to alter the property, and, optionally,

iii) testing the resulting Termamyl-like .alpha.-amylase variant with respect to the property in question.

5 Claims, 11 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 13

----- KWIC -----

Other Reference Publication - OREF (5):

Gray et al., "Structural <u>Genes Encoding The Thermophilic .alpha.-Amylases</u> of Bacillus Stearothermophilus And Bacillus Licheniformis", Journal of Bacteriology, vol. 166, No. 2, May 1996, pp. 635-643.

5981243

DOCUMENT-IDENTIFIER: US 5981243 A

TITLE:

Purified myceliophthora laccases and nucleic acids

encoding same

DATE-ISSUED:

November 9, 1999

INVENTOR-INFORMATION:

CITY NAME Davis Berka; Randy Michael

ZIP CODE COUNTRY STATE 95616 N/A CA

Brown; Stephen H.

Davis

CA 95616 N/A

Xu; Feng Schneider; Palle Woodland DK-2750 Ballerup

95776 N/A N/A N/A

DK Oxenb.o slashed.ll; Karen M. DK-2920 Charlottenlund N/A DK N/A

Aaslyng; Dorrit A.

Gartnerkrogen 69

N/A

N/A

DK

APPL-NO:

08/ 939218

DATE FILED: September 29, 1997

PARENT-CASE:

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a divisional of U.S. application Ser. No. 08/441,146 filed May 15, 1995, now abandoned, which a continuation-in-part of application Ser. No. 08/253,781 filed Jun. 3, 1994, now abandoned, the contents of which are fully incorporated herein by reference.

US-CL-CURRENT: 435/189, 536/23.2, 8/401

ABSTRACT:

The present invention relates to isolated nucleic acid constructs containing a sequence encoding a Myceliophthora laccase, and the laccase proteins encoded thereby.

18 Claims, 3 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 6

----- KWIC -----

Detailed Description Text - DETX (55):

The construction strategy for the laccase expression vector pRaMB5 is outlined in FIG. 3. The promoter directing transcription of the laccase gene is obtained from the A. oryzae .alpha.-amylase (TAKA-amylase) gene (Christensen et al., supra), as well as the TAKA-amylase terminator region. The plasmid is constructed first by modifying pMWR3 by inserting a small linker which contains an Apal site between the Swal and Nsil sites, creating a plasmid called pMWR3-SAN. Pful polymerase-directed PCR (Stratagene, La Jolla, Calif.) is used to amplify a short DNA segment encoding the 5'-portion of MtL, from the start codon to an internal Pstl site (approximately 0.5 kb). The forward primer for this PCR reaction is designed to create an EcoRI site just upstream of the start codon. Next, the amplified fragment is digested with EcoRI and Pstl[during this step, the EcoRl site is made blunt by treatment with dNTPs and DNA polymerase I(Klenow fragment)] and purified by agarose gel electrophoresis. The 3' portion of the M. thermophila coding region is excised from pRaMB2 as a 2kb Pstl-Apal fragment(this segment also contains approximately 110 bp from the 3'-untranslated region). These two fragments are combined with Swal- and Apal-cleaved pMWR3-SAN in a three-part ligation reaction to generate the laccase expression vector pRaMB5.

5958739

DOCUMENT-IDENTIFIER: US 5958739 A

TITLE:

Mutant .alpha.-amylase

DATE-ISSUED:

September 28, 1999

INVENTOR-INFORMATION:

INVENTOR-INFORM	IATION.				
NAME	CITY	STATE	ZIP C	ODE CO	JNTRY
Mitchinson; Colin	Palo Alto	CA	N/A	N/A	
Requadt; Carol	Palo Alto	CA	N/A	N/A	
Ropp; Traci	Palo Alto	CA	N/A	N/A	
Solheim; Leif P.	Palo Alto	CA	N/A	N/A	
Ringer; Christopher	Palo Alto	CA	N/A	N/A	
Day; Anthony	Palo Alto	CA	N/A	N/A	

APPL-NO:

08/704706

DATE FILED: February 20, 1997

PCT-DATA:

APPL-NO: PCT/US96/09089 DATE-FILED: June 6, 1996 WO96/39528 PUB-NO: PUB-DATE: Dec 19, 1996 371-DATE: Feb 20, 1997 102(E)-DATE:Feb 20, 1997

US-CL-CURRENT: 435/99, 435/201, 435/202, 435/203, 435/204, 435/252.3

. 435/252.31 , 435/254.11 , 435/320.1 , 435/325 , 435/410 , 510/226 , 510/300 , 510/305 , 510/320 , 510/374 , 510/392

, 536/23.2

ABSTRACT:

Novel .alpha.-amylase enzymes are disclosed in which one or more asparagine residues are substituted with a different amino acid or deleted. The disclosed .alpha.-amylase enzymes show altered or improved low pH starch hydrolysis performance, stability and activity profiles.

32 Claims, 13 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 17

----- KWIC -----

Other Reference Publication - OREF (26):
Gray et al., "Structural **Genes Encoding the Thermophilic .alpha.-amylases** of Bacillus stearothermophilus and Bacillus licheniformis," J Bacteriol (1986)

166:635-643.

5849549

DOCUMENT-IDENTIFIER: US 5849549 A

TITLE:

Oxidatively stable alpha-amylase

DATE-ISSUED:

December 15, 1998

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE COUNTRY

Barnett; Christopher C.

South San Franciso Clinton

CA N/A

Solheim; Leif P. Mitchinson: Colin IΑ

CA

N/A N/A N/A

N/A

N/A

Power: Scott D. Requadt; Carol A. Half Moon Bay San Bruno

CA CA N/A

N/A

Tiburon

N/A N/A

APPL-NO:

08/468698

DATE FILED: June 6, 1995

PARENT-CASE:

RELATED APPLICATION

This is a divisional of U.S. Ser. No. 08/194,664 filed Feb. 10, 1994, now pending, which is a continuation-in-part of U.S. Ser. No. 08/016,395 filed Feb. 11, 1993, abandoned.

US-CL-CURRENT: 435/99, 435/202, 536/23.2

ABSTRACT:

Novel alpha-amylase mutants derived from the DNA sequences of naturally occurring or recombinant alpha-amylases are disclosed. The mutant alpha-amylases, in general, are obtained by in vitro modifications of a precursor DNA sequence encoding the naturally occurring or recombinant alpha-amylase to generate the substitution (replacement) or deletion of one or more oxidizable amino acid residues in the amino acid sequence of a precursor alpha-amylase. Such mutant alpha-amylases have altered oxidative stability and/or altered pH performance profiles and/or altered thermal stability as compared to the precursor. Also disclosed are detergent and starch liquefaction compositions comprising the mutant amylases, as well as methods of using the mutant amylases.

2 Claims, 28 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 22

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 KWIC	

Other Reference Publication - OREF (5):

Gray, et al., "Structural Genes Encoding the Thermophilic .alpha.-amylases of Bacillus sterothermophilus and Bacillus licheniformis" J. Bact. 166(2):635-643 (May 1986).

5840851

DOCUMENT-IDENTIFIER: US 5840851 A

TITLE:

Purification of hemoglobin

DATE-ISSUED:

November 24, 1998

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Plomer; J. Jeffrey	Broomfield	CO	80020	N/A
Ryland; James R.	Louisville	CO	80027	N/A
Matthews; Maura-Ann	H. Boulder	CC	8030	4 N/A
Traylor; David W.	Wheat Ridge	CO	80033	N/A
Milne; Erin E.	Broomfield	CO	80020	V/A
Durfee; Steven L.	Denver	CO	80207	N/A
Mathews; Antony J.	Louisville	CO	80027	N/A
Neway; Justinian O.	Longmont	CO	80503	N/A

APPL-NO:

08/438511

DATE FILED: May 10, 1995

PARENT-CASE:

CROSS-REFERENCES TO RELATED APPLICATIONS

This application is a continuation-in-part of application Ser. No. 08/339,304, filed Nov. 14, 1994, now abandoned which is a continuation-in-part of application Ser. No. Ser. No. 08/097,273, filed Jul. 23, 1993, now abandoned, both incorporated herein by reference in their entirety.

US-CL-CURRENT: 530/385, 530/412, 530/413, 530/416

ABSTRACT:

The present invention generally relates to methods for purifying hemoglobin solutions and to hemoglobin solutions obtained by the methods. In one aspect, such methods include removing contaminants in crude hemoglobin-containing lysates with heat treatment. In a further aspect, the present invention provides methods for producing substantially purified hemoglobin solutions using immobilized metal affinity chromatography, optionally following by anion exchange chromatography.

51 Claims, 2 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 2

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 L/ AA IC >	

Other Reference Publication - OREF (34):
Tsukagoshi, N. et al/Cloning and Expression of a <u>Thermophilic</u>
<u>.alpha.-Amylase Gene</u> from Bacillus Stearothermophilus in Escherichia Coli/Mol.
Gen Genet/(1984)/193: 58-63.

5824532

DOCUMENT-IDENTIFIER: US 5824532 A

TITLE:

Oxidativley stable alpha-amylase

DATE-ISSUED:

October 20, 1998

INVENTOR-INFORMATION:

ZIP CODE COUNTRY NAME STATE South San Francisco Barnett; Christopher C. CA N/A N/A Mitchinson: Colin Half Moon Bay N/A CA N/A Power: Scott D. San Bruno CA N/A N/A Requadt; Carol A. Tiburon CA N/A N/A

APPL-NO:

08/468220

DATE FILED: June 6, 1995

PARENT-CASE:

RELATED APPLICATIONS

This is a divisional of U.S. Ser. No. 08/194,664 filed Feb. 10, 1994, now pending which is a continuation-in-part of U.S. Ser. No. 08/016,395 filed Feb. 11, 1993 now abandoned.

US-CL-CURRENT: 435/202, 435/201, 435/203, 435/204, 435/252.3, 435/252.31 , 435/320.1 , 435/71.2 , 536/23.2 , 536/23.7

ABSTRACT:

Novel alpha-amylase mutants derived from the DNA sequences of naturally occurring or recombinant alpha-amylases are disclosed. The mutant alpha-amylases, in general, are obtained by in vitro modifications of a precursor DNA sequence encoding the naturally occurring or recombinant alpha-amylase to generate the substitution (replacement) or deletion of one or more oxidizable amino acid residues in the amino acid sequence of a precursor alpha-amylase. Such mutant alpha-amylases have altered oxidative stability and/or altered pH performance profiles and/or altered thermal stability as compared to the precursor. Also disclosed are detergent and starch liquefaction compositions comprising the mutant amylases, as well as methods of using the mutant amylases.

11 Claims, 28 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 22

KWIC	
 TANK.	

Other Reference Publication - OREF (2):

G. Gray et al., "Structural <u>Gene Encoding the Thermophillic .alpha.-Amylases</u> of Bacillus stearothermophilus and Bacillus licheniformis", J. Bact. 166(2) 635-643, (May 1986).

5756714

DOCUMENT-IDENTIFIER: US 5756714 A

TITLE:

Method for liquefying starch

DATE-ISSUED:

May 26, 1998

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE COUNTRY

Antrim: Richard L.

Solon

N/A N/A

Mitchinson; Colin

Half Moon Bay

CA N/A N/A

Solheim; Leif P.

Clinton

IA N/A N/A

IA

APPL-NO:

08/411038

DATE FILED: March 27, 1995

PARENT-CASE:

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation-in-part of U.S. application Ser. No. 08/401,325 filed Mar. 9, 1995, now abandoned and which is incorporated herein by reference in its entirety.

US-CL-CURRENT: 536/102, 435/202, 435/203, 435/204, 435/205, 435/96 , 435/99

ABSTRACT:

According to the invention a method is provided for liquefying starch comprising the steps of treating the starch prior to or simultaneously with liquefying the starch to inactivate and/or remove the enzyme inhibiting composition present in the starch and form treated starch; adding .alpha.-amylase to the treated starch; and reacting the treated starch for a time and at a temperature effective to liquefy the treated starch. Effective means to treat the starch include the addition of a phytate degrading enzyme and heat treatment, optionally followed by filtration or centrifugation, of granular starch or a starch solution.

17 Claims, 0 Drawing figures	
Exemplary Claim Number:	1

----- KWIC -----

Other Reference Publication - OREF (8):

Gray, et al., "Structural <u>Genes Encoding the Thermophilic .alpha.-Amylases</u> of Bacillus sterothermophilus and Bacillus lichenmiformis" J. of Bacteriology 166(2):635-643 (May 1986).

5707841

DOCUMENT-IDENTIFIER: US 5707841 A

TITLE:

Process of producing highly transformable bacterial

cells and cells produced thereby

DATE-ISSUED:

January 13, 1998

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE COUNTRY

Greener; Alan L.

San Diego

CA N/A

N/A

APPL-NO:

08/637003

DATE FILED: April 18, 1996

PARENT-CASE:

This is a continuation of application Ser. No. 08,151,577, filed Nov. 12, 1993, now U.S. Pat. No. 5,512,468.

US-CL-CURRENT: 435/488, 435/252.33, 435/252.8

ABSTRACT:

The invention provided herein includes gram negative bacteria cells containing a gene encoding an enzyme with carbohydrate degrading activity that had been rendered competent to transformation. Carbohydrate degrading enzymes of interest for use in the invention include alpha-amylase. The competent cells of the subject invention may be frozen so as to provide for prolonged storage. Other aspects of the invention include methods for rendering gram negative bacterial cells, such as E. coli cells competent to transformation. These methods involve the step of transferring a gene encoding an enzyme with carbohydrate degrading activity into E. coli cells and subsequently rendering the cells competent using any of a variety of competency inducing procedures.

13 Claims, 0 Drawing figures

Exemplary Claim Number:

----- KWIC -----

Brief Summary Text - BSTX (14):

The term carbohydrate-degrading enzyme as used herein refers to enzymes capable of hydrolyzing at least one type of linkage present between the constituent monosaccharide units of a carbohydrate molecule. The term "starch-degrading enzyme" as used herein refers to enzymes capable of

hydrolyzing at least one type of linkage present between the constituent monosaccharide units of a starch molecule. The term "alpha-amylase" as used herein refers to enzymes capable of catalyzing hydrolysis of the .alpha.-1.fwdarw.4 glucosidic linkages of polysaccharides containing such glucosidic linkages such as starch or glycogen. Preferred carbohydrate degrading enzymes are starch degrading enzymes. Preferred starch degrading enzymes are alpha-amylases. Particularly preferred starch degrading enzymes for use in the invention are alpha-amylases. Particularly preferred starch degrading enzymes for use in the invention are alpha-amylase from a recently isolated uncharaterized thermophilic bacterium. The polynucleotide sequence encoding this alpha-amylase from the uncharacterized thermophilic bacterium can be found on the FAMY plasmid present in the E. coli strains having the ATCC accession numbers 69480, 69481, and 69482.

Brief Summary Text - BSTX (18):

The introduction of a genetic construction for the expression of a carbohydrate degrading enzyme into E. coli serves to increase the transformation efficiency of compositions of E. coli cells rendered competent for a wide variety of E. coli strains. The genotype of an E. coli cell strain containing a genetic construction for the expression of a carbohydrate degrading enzyme may be selected so as to be particularly useful for a given genetic engineering experiment. For example, cloning vectors that are screenable because of LacZ, alpha, fragment complementation may contain a particular mutation within the LacZ gene. Similarly, the cell may contain various other deletions or mutations in order to provide for complementation by the transforming DNA. The host cell may either possess or lack a restriction-modification system in order to expedite cloning. The host cells may also lack one or more recombination systems, e.g., RecA, RecBC. Preferred E. coli strains for use in the invention are cells that contain a mutation in the deoR gene, as described in Hanahan, U.S. Pat. No. 4,851,348. Particularly preferred strains of E. coli for use in the invention are the XL1-Blue.TM. strain (Stratagene, La Jolla, Calif.), the XL1-Blue MR strain. and the SURE.TM. strain (Stratagene, La Jolla, Calif.) that have been modified by the addition of a genetic construction for the expression of alpha-amylase isolated from a thermophilic bacteria and have the ATCC accession numbers 69480, 69481 and 69482, respectively. The plasmid containing the alpha-amylase gene in the E. coli strains having ATCC accession numbers 69480, 69481 and 69482 may be readily transferred to other strains of bacteria using techniques well known to the person of average skill in the art. Similarly, the person of average skill in the art may excise the alpha amylase gene from plasmids in the E. coli strains having accession numbers 69480, 69481 and 69482 and transfer the alpha amylase gene to a new genetic construct prior to transferring the gene to a new strain of bacteria.

Detailed Description Text - DETX (6):

An <u>alpha-amylase gene</u> from a recently isolated uncharacterized <u>thermophilic</u> bacterium was initially inserted onto an autonomously replicating plasmid DNA element, and then was introduced into several E. coli strains through conjugation. This <u>alpha-amylase gene</u> can be found in the plasmids present in the E. coli strains having the ATCC accession numbers 69480, 69481, and 69482.

The resultant <u>alpha-amylase gene</u> containing strains and the respective parent strain were rendered competent using the procedure of Hanahan (J. Mol. Biol. (1983)). The transformation efficiency of the <u>alpha-amylase gene containing strains were compared with the transformation of similar strains lacking the alpha-amylase gene.</u>

Detailed Description Text - DETX (11):

Cloning <u>alpha-amylase gene</u> onto RSF1010 derivative pAL205. pBM100, a pBluescript.TM. II vector derivative containing the amylase gene from a <u>thermophilic</u> bacterium was digested with Xbal and HindIII and the appropriate DNA fragment was isolated and ligated to pAL205 digested with these same two enzymes. The ligation mix was transformed into SCS 1 and Tet.sup.R Cam.sup.R colonies selected. Plasmid DNA from these transformants was isolated and subjected to restriction enzyme analysis to confirm the presence of the amylase gene.

Claims Text - CLTX (4):

2. The method of claim 1, wherein the <u>alpha-amylase gene</u> is isolated from a **thermophilic** bacterium.

Claims Text - CLTX (15):

10. The competent bacterial cell according to claim 9, wherein the alpha-amylase gene is isolated from a thermophilic bacterium.

Other Reference Publication - OREF (4):

Tsukagoshi et al., 1984, "Cloning and Expression of a <u>Thermophilic</u> <u>.alpha.-Amylase Gene</u> from Bacillus stearothermophilus in Escherichia coli," Mol. Gen. Genet. 193:58-63.

DOCUMENT-IDENTIFIER: US 5665869 A

TITLE:

Method for the rapid removal of protoporphyrin from protoporphyrin IX-containing solutions of hemoglobin

CA

N/A

N/A

DATE-ISSUED: September 9, 1997

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Ryland; James R. Louisville CO N/A N/A Matthews; Maura-Ann H. Boulder CO N/A N/A Ernst; Ulrich P. Lafayette CO N/A N/A Houk; Daniel E. Concord CA N/A N/A Traylor; David W. Wheat Ridge CO N/A N/A Williams; Lee R. Concord

APPL-NO: 08/153071

DATE FILED: November 15, 1993

US-CL-CURRENT: 530/412, 530/385

ABSTRACT:

The present invention relates to a method for the production of a substantially protoporphyrin IX free hemoglobin solution comprising: rapidly heating a crude protoporphyrin IX-containing hemoglobin solution for a relatively short time and at a relatively high temperature to reduce protoporphyrin IX-containing hemoglobin to insignificant levels in said protoporphyrin IX-containing hemoglobin solution.

38 Claims, 12 Drawing figures

Exemplary Claim Number: 38

Number of Drawing Sheets: 12

----- KWIC -----

Other Reference Publication - OREF (18):

Tsukagoshi, N. et al/Cloning and Expression of a Thermophilic <u>-alpha.-Amylase Gene</u> from Bacilus stearothermophilus in Escherichia coli/Mol. Gen. Genet./(1984)/193, 58-63.

5512468

DOCUMENT-IDENTIFIER: US 5512468 A **See image for Certificate of Correction**

TITLE:

Process of producing highly transformable bacterial

cells and cells produced thereby

DATE-ISSUED:

April 30, 1996

INVENTOR-INFORMATION:

NAME

STATE ZIP CODE COUNTRY

Greener; Alan L.

San Diego

CA N/A N/A

APPL-NO:

08/ 151577

DATE FILED: November 22, 1993

US-CL-CURRENT: 435/488, 435/252.33, 435/252.8

ABSTRACT:

The invention provided herein includes gram negative bacteria cells containing a gene encoding an enzyme with carbohydrate degrading activity that had been rendered competent to transformation. Carbohydrate degrading enzymes of interest for use in the invention include alpha-amylase. The competent cells of the subject invention may be frozen so as to provide for prolonged storage.

Other aspects of the invention include methods for rendering gram negative bacterial cells, such as E. coli cells competent to transformation. These methods involve the step of transferring a gene encoding an enzyme with carbohydrate degrading activity into E. coli cells and subsequently rendering the cells competent using any of a variety of competency inducing procedures.

11 Claims, 0 Drawing figures

Exemplary Claim Number:

----- KWIC -----

Brief Summary Text - BSTX (14):

The term "starch-degrading enzyme" as used herein refers to enzymes capable of hydrolyzing at least one type of linkage present between the constituent monosaccharide units of a carbohydrate molecule. The term "starch-degrading enzyme" as used herein refers to enzymes capable of hydrolyzing at least one type of linkage present between the constituent monosaccharide units of a starch molecule. The term "alpha-amylase" as used herein refers to enzymes

capable of catalyzing hydrolysis of the .alpha.-1.fwdarw.4 glucosidic linkages of polysaccharides containing such glucosidic linkages such as starch or glycogen. Preferred carbohydrate degrading enzymes are starch degrading enzymes. Preferred starch degrading enzymes are alpha-amylases. Particularly preferred starch degrading enzymes for use in the invention are alpha-amylases. Particularly preferred starch degrading enzymes for use in the invention are alpha-amylases isolated from thermophilic bacteria, especially the alpha-amylase gene from a recently isolated uncharaterized thermophilic bacterium. The polynucleotide sequence encoding this alpha-amylase from the uncharacterized thermophilic bacterium can be found on the FAMY plasmid present in the E. coli strains having the ATCC accession numbers 69480, 69481, and 69482.

Brief Summary Text - BSTX (18):

The introduction of a genetic construction for the expression of a carbohydrate degrading enzyme into E. coli serves to increase the transformation efficiency of compositions of E. coli cells rendered competent for a wide variety of E. coli strains. The genotype of an E. coli cell strain containing a genetic construction for the expression of a carbohydrate degrading enzyme may be selected so as to be particularly useful for a given genetic engineering experiment. For example, cloning vectors that are screenable because of LacZ.alpha. fragment complementation may contain a particular mutation within the LacZ gene. Similarly, the cell may contain various other deletions or mutations in order to provide for complementation by the transforming DNA. The host cell may either possess or lack a restriction-modification system in order to expedite cloning. The host cells may also lack one or more recombination systems, e.g., RecA, RecBC. Preferred E. coli strains for use in the invention are cells that contain a mutation in the deoR gene, as described in Hanahan, U.S. Pat. No. 4,851,348. Particularly preferred strains of E. coli for use in the invention are the XL1-Blue.TM. strain (Stratagene, La Jolla, Calif.), the XL1-Blue MR strain, and the SURE.TM. strain (Stratagene, La Jolla, Calif.) that have been modified by the addition of a genetic construction for the expression of alpha-amylase isolated from a thermophilic bacteria and have the ATCC accession numbers 69480, 69481 and 69482, respectively. The plasmid containing the alphaamylase gene in the E. coli strains having ATCC accession numbers 69480, 69481 and 69482 may be readily transferred to other strains of bacteria using techniques well known to the person of average skill in the art. Similarly, the person of average skill in the art may excise the alpha amylase gene from plasmids in the E. coli strains having accession numbers 69480, 69481 and 69482 and transfer the alpha amylase gene to a new genetic construct prior to transferring the gene to a new strain of bacteria.

Detailed Description Text - DETX (5):

An <u>alpha-amylase gene</u> from a recently isolated uncharacterized <u>thermophilic</u> bacterium was initially inserted onto an autonomously replicating plasmid DNA element, and then was introduced into several E. coli strains through conjugation. This <u>alpha-amylase gene</u> can be found in the plasmids present in the E. coli strains having the ATCC accession numbers 69480, 69481, and 69482. The resultant <u>alpha-amylase gene</u> containing strains and the respective parent strain were rendered competent using the procedure of Hanahan (J. Mol. Biol. (1983)). The transformation efficiency of the <u>alpha-amylase gene containing</u>

strains were compared with the transformation of similar strains lacking the alpha-amylase gene.

5366883

DOCUMENT-IDENTIFIER: US 5366883 A

TITLE:

.alpha.-amylase gene

DATE-ISSUED:

November 22, 1994

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP C	DDE CC	UNTRY
Asada; Kiyozo	Shiga	N/A	N/A	JP	
Uemori; Takashi	Shiga	N/A	N/A	JP	
Mukai; Hiroyuki	Shiga	N/A	N/A	JP	
Kato; Ikunoshin	Kyoto	N/A	N/A	JP	
Laderman; Kenneth	Baltimore	ME) N/.	A N/	Α
Anfinsen; Christian B.	Baltimore	MD	N/A	N/A	

APPL-NO:

07/894212

DATE FILED: June 9, 1992

US-CL-CURRENT: 435/202, 435/252.3 , 435/252.31 , 435/252.33 , 435/320.1

, 435/69.1 , 435/71.2 , 536/23.1 , 536/23.2

ABSTRACT:

The present invention relates, in general, to a cloned .alpha.-amylase gene. and, in particular, to a cloned hyperthermophilic .alpha.-amylase gene and to methods of producing .alpha.-amylase using same.

13 Claims, 9 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 9

----- KWIC -----

Abstract Text - ABTX (1):

The present invention relates, in general, to a cloned alpha.-amylase gene, and, in particular, to a cloned hyperthermophilic .alpha.-amylase gene and to methods of producing .alpha.-amylase using same.

Brief Summary Text - BSTX (2):

The present invention relates, in general, to a cloned .alpha.-amylase gene, and, in particular, to a cloned hyperthermophilic .alpha.-amylase gene and to

methods of producing .alpha.-amylase using same.

Brief Summary Text - BSTX (7):

Using genetic engineering technology, it is theoretically possible to clone genes and produce the enzymes that they encode in quantities sufficient for industrial application. A number of **genes coding for thermophilic .alpha.-amylases** have been isolated and subsequently expressed in E. coli and B. subtilis (Fukusumi et al, Eur. J. Biochem., 98: 95 (1985), Tsukagoshi et al, Mol. Gen. Genet., 195: 58 (1984), Tsukagoshi et al, J. Bacteriology, 164: 1182 (1985)). The temperature at which the genes are endogenously translated does not seem to have an effect on the expression in transformation competent cells. Thus it is possible to produce **thermophilic** enzymes in host cells grown at ambient temperature. However, no **genes coding for hyperthermophilic .alpha.-amylases** have ever been successfully cloned.

Brief Summary Text - BSTX (9):

The present invention provides, for the first, time a cloned **sequence encoding a hyperthermophilic .alpha.-amylase**. The availability of this sequence makes possible the industrial scale production of this enzyme.

Brief Summary Text - BSTX (11):

The present invention relates to an isolated DNA segment having a nucleotide **sequence that encodes .alpha.-amylase**, specifically, a **hyperthermophilic** .alpha.-amylase. The invention further relates to a recombinant method of producing **hyperthermophilic** .alpha.-amylase. The invention also relates to an expression vector suitable for use in such a method.

Brief Summary Text - BSTX (12):

It is a general object of the invention to provide a gene encoding a hyperthermophilic .alpha.-amylase.

Detailed Description Text - DETX (6):

The Examples that follow make reference to the hyperthermophilic bacteria, P. furiosus (deposited at Deutsche Sammlung von Mikroorganismen with the identification number of DSM 3638). The procedures for cloning the alpha.-amylase gene from this bacteria and for preparing transformants carrying the gene, can be described in general terms as follows:

5364782

DOCUMENT-IDENTIFIER: US 5364782 A

TITLE:

Mutant microbial .alpha.-amylases with increased

thermal, acid and/or alkaline stability

DATE-ISSUED:

November 15, 1994

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Quax: Wilhelmus J.

Voorschoten Brussels

Naaldwijk

N/A N/A NL BE

Laroche; Yves Vollebregt; Adrianus W. H. N/A N/A N/A

N/A NL

BE

Stanssens; Patrick

St. Denijs Westrem

N/A N/A

BE

Lauwereys; Marc

Haaltert

N/A N/A

APPL-NO:

07/623953

DATE FILED: November 29, 1990

FOREIGN-APPL-PRIORITY-DATA:

COUNTRY

APPL-NO

APPL-DATE

EΡ

89201735

June 29, 1989

PCT-DATA:

APPL-NO: PCT/EP90/01042 DATE-FILED: June 27, 1990 WO91/00353 PUB-NO: PUB-DATE: Jan 10, 1991 371-DATE: Dec 2, 1990

102(E)-DATE:Dec 2, 1990

US-CL-CURRENT: 435/202, 435/252.3, 435/263, 435/275, 435/320.1, 536/23.2

ABSTRACT:

Thermostable and acid stable .alpha.-amylases are provided as expression products of genetically engineered .alpha.-amylase genes isolated from microorganisms, preferably belonging to the class of Bacilli. Both chemical and enzymatic mutagenesis methods are e.g. the bisulphite method and enzymatic misincorporation on gapped heteroduplex DNA. The mutant .alpha.-amylases have superior properties, e.g. improved thermostability over a broad pH range, for industrial application in starch processing and textile desizing.

6 Claims, 15 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 24

Other Reference Publication - OREF (1):
Gray et al., Structural <u>Genes Encoding the Thermophilic alpha-Amylases</u> of Bacillus stearothermophilus and Bacillus licheniformis, J. Bacteriol. (1986) 166:635-643.

4946789

DOCUMENT-IDENTIFIER: US 4946789 A

TITLE:

Bacillus brevis strains and application thereof

DATE-ISSUED:

August 7, 1990

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE COUNTRY

Udaka; Shigezo

Aichi Chiba

N/A N/A JΡ

Takagi; Hiroaki Kadowaki; Kiyoshi

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APPL-NO:

07/043459

DATE FILED: April 28, 1987

FOREIGN-APPL-PRIORITY-DATA:

COUNTRY

APPL-NO

APPL-DATE

JP

61-198120

August 26, 1986

US-CL-CURRENT: 435/252.3, 435/252.31, 435/69.1, 435/71.1, 435/71.2

ABSTRACT:

Bacillus brevis strains which produce a large amount of protein but no protease out of the cells are disclosed. These strains are highly useful as hosts in genetic engineering.

2 Claims, 4 Drawing figures

Exemplary Claim Number:

Number of Drawing Sheets: 2

----- KWIC -----

Detailed Description Text - DETX (24):

Cloning of Thermophilic .alpha.-amylase Gene into E. Coli

Detailed Description Text - DETX (27):

SUBCOLONING OF THERMOPHILIC .alpha.-AMYLASE GENE

4493893

DOCUMENT-IDENTIFIER: US 4493893 A

TITLE:

Process for cloning the gene coding for a thermostable alpha-amylase into Escherichia coli and Bacillus subtilis

DATE-ISSUED:

January 15, 1985

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE COUNTRY

Mielenz; Jonathan R. Mickel; Susan

LaGrange Park

IL N/A N/A

N/A

LaGrange Park

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APPL-NO:

06/472646

DATE FILED: March 11, 1983

PARENT-CASE:

This application is a continuation of application Ser. No. 225,287 filed Jan. 15, 1981, now abandoned.

US-CL-CURRENT: 435/91.41, 435/201, 435/320.1, 435/69.1, 435/69.2

ABSTRACT:

An improved process for producing a thermostable alpha-amylase enzyme is described. The gene coding for the alpha-amylase is incorporated into a chimeric plasmid which is produced in multiple copies by a host microorganism.

22 Claims, 0 Drawing figures

Exemplary Claim Number:

----- KWIC -----

Brief Summary Text - BSTX (22):

The chimeric plasmids of this invention are prepared using DNA from a naturally-occurring donor microorganism which contains a gene coding for a thermostable alpha-amylase enzyme. Suitable donor microorganisms are found in the thermophilic bacteria classified as Bacillus stearothermophilus (abbreviated B. stearothermophilus) and Thermus flavus (abbreviated T. flavus). Strains of Bacillus licheniformis (abbreviated B. licheniformis) are also suitable donor microorganisms. Strains of B. stearothermophilus particularly suitable for the use as a source of donor DNA are those strains selected from the group consisting of B. stearothermophilus, ATCC Nos. 31,195; 31,196; 31,197; 31,198; 31,199 and 31,783, variants and mutants thereof and submutants

of said mutants.